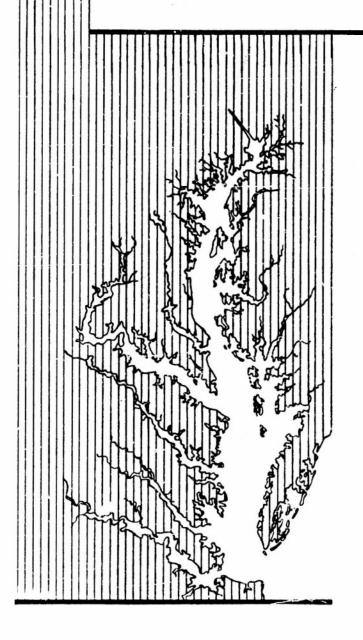


THE CHESAPEAKE BAY INSTITUTE of The Johns Hopkins University



TECHNICAL REPORT VII

A STUDY OF FLUSHING IN THE DELAWARE MODEL

by D.W. Pritchard

Reference 54-4 April 1954 THIS REPORT HAS BEEN DELIMITED AND CLEARED FOR PUBLIC RELEASE UNDER DOD DIRECTIVE 5200.20 AND NO RESTRICTIONS ARE IMPOSED UPON ITS USE AND DISCLOSURE.

DISTRIBUTION STATEMENT A

APPROVED FOR PUBLIC RELEASE; DISTRIBUTION UNLIMITED,

CHESAPEAKE BAY INSTITUTE THE JOHNS HOPKINS UNIVERSITY

TECHNICAL REPORT VII

A STUDY OF FLUSHING IN THE DELAWARE MODEL

by

D. W. Pritchard

This report contains results of work carried out for: the Office of Naval Research of the Department of the Navy under research project NR 083-070, Contract Nonr 248(30); and research project NR 084-005, Contract Nonr 248(07); and for the U.S. Navy Hydrographic Office.

This report does not necessarily constitute final publication of the material presented.

Reference 54-4 April 1954 D. W. Pritchard Director

A STUDY OF FLUSHING IN THE DELAWARE MODEL

Introduction

Late in 1951 the Chesapeake Bay Institute was asked to assist the Oceanographic Division of the Hydrographic Office in a study of the flushing problem, using the Delaware River and Bay Model located at the U. S. Army Engineers Waterways Experiment Station, Vicksburg, Mississippi. The first test made showed that the dye used as a tracer contaminant was adsorbed in large quantities by the concrete bottom of the model, the glass sample bottles, and even by the dust which settled from the air into the model. Further tests were delayed until February 1952 while the model was painted with adsorption inhibiting plastic paint.

Eight separate tests were made on the model by introducing a known volume of dye and tracing the changes in the distribution of the dye by photometric analysis of samples. Considerable reduction of data was necessary to compensate for the adsorption and the body of this report will deal only with the reduced data.

Several appendices have been included. Appendix I deals with the sampling gear, the method of sampling, and the methods of field analysis. Tables of observed dye concentrations, by station and tidal cycle are given in Appendix II. Appendix III gives the reduction methods used to compensate for the adsorption and tables of corrected concentrations by station and tidal cycle. Appendix IV contains auxiliary data on salinity, high and low water profiles, and fresh water inflow as well as tables of

cross-sectional areas, intersectional volumes, and accumulated volumes for each 5000 foot station.

Description of Flushing Tests

The Delaware Model

The region represented by the Delaware model includes all of the Delaware Bay and the tidal section of the Delaware River, from the Capes to Trenton. The model is constructed of concrete on a horizontal scale of 1 to 1000 and a vertical scale of 1 to 100. A chart of the model is given in Figure 1.

Information about the hydraulic adjustment and verification, and the salinity verification are presented in various reports of the Water-ways Experiment Station (Corps of Engineers, U. S. Army 1951(a), 1951(b), and 1952). These reports were sufficiently conclusive to suggest that valuable information on flushing in the Delaware River and Bay could be obtained from the model.

The Flushing Tests

In the following discussion the tests are grouped according to the point of release. In Tests 1-A, 1, and 4 the dye was introduced at Station 52.5 which is located a short distance below Philadelphia. Station 292.5 below Artificial Island, was used as the release point in Tests 2 and 3. The dye was released from Station 111 located near Chester, for Tests 5, 6, and 7.

While some treatment is presented of the lateral and vertical distributions, the main discussion here will center on the variation in concentration along the axis of the estuary. In each case the data represent conditions at the time of high water slack.

The manner of introducing the dye is discussed in detail in Appendix I. Briefly, the dye was introduced at high water slack in a cylinder constructed so that the sides could be freed from the bottom and pulled out of the water, leaving the dye suspended momentarily in the water in a cylindrical shape. Then as the ebb tidal movement started, the dye was rapidly spread in both a lateral and a longitudinal direction. The diameter of the cylindrical container was 0.74 ft, equivalent to 740 ft in the prototype. The height of dye in the container was adjusted to be even with the surface of the water in the model at the time of release.

Test 1-A, 1, and 4 - Release Point 52.5

Test 1-A, which was the trial run made in December 1951, and Test 1, which was the first of the series of runs made in February 1952, were duplicate tests, the dye being released at Station 52.5 under conditions of mean tide and mean river flow (12,350 sec ft at Trenton). The dye concentration at release was 1000 ppm, and even though at this location in the tidal river the water is practically fresh, no density effects were observed.

Upon release the dye cylinder moved seaward with the ebbing tide while mixing laterally and longitudinally. After one tidal cycle the dye had spread longitudinally over nearly one tidal excursion assuming a nearly normal distribution, and after two tidal cycles it had a nearly

uniform distribution laterally.

It was observed that in pockets in the shore line small eddies would sometimes trap water containing a high concentration of dye.

After the high concentration of dye had passed, the entrapped dye would be slowly fed back into the main stream. This phenomenon was observed both on the ebb and on the flood tide and probably materially assists the spread of the highly concentrated contaminant. Figure 2 is a schematic representation of the process.

Figures 3a and 3b show the longitudinal distributions of the contaminant concentration from 1 to 58 tidal cycles after the dye was released for Test 1-A. (Test 1 was substantially the same as Test 1-A.) The solid vertical column shown at Station 52.5 represents the initial dye volume. After one tidal cycle the contaminant had spread over nearly one tidal excursion and the distribution approximated the normal curve. With increasing time the spread increased while the peak concentration moved seaward.

The rate of decrease of the peak concentration seems to be exponential and Figure 4 shows a semilog plot of peak concentration against time in tidal cycles for both Test 1-A and Test 1. Two exponential decay rates are evident. For the first two tidal cycles the rate of decrease is extremely rapid, falling from 1000 ppm for time zero to less than 6 ppm after two tidal cycles. For times greater than two tidal cycles, decay progresses at a much slower rate. It would seem, then, that the process governing the dispersion of the contaminant during the first two

tidal cycles are fundamentally different from those acting after the second tidal cycle.

Figure 5 shows the movement of the peak concentration with time in tidal cycles by the solid line. The dashed line is the calculated net downstream water movement based on river inflow. It is to be noted that the peak concentration moves downstream more slowly than the net downstream water movement. This was the case in every test.

If instead of following the movement of the peak concentration, we consider the concentration of dye at a fixed point, we get a different picture. Figure 6 shows the change in concentration with time for stations above the release point while Figure 7 does the same thing for stations below the release point. For stations well above the release point (e.g. 20 and 30) an increase in concentration occurs during the first few tidal cycles and is followed by an approximately exponential decay after the local peak has been passed. For upstream stations nearer the release point (e.g. 40 and 50) the initial rise occurs during the first tidal cycle. The decay is very rapid during the first few tidal cycles becoming approximately exponential as time passes. For stations downstream from the release point the decay becomes approximately exponential with the passage of time but the initial increase prior to the passage of the peak concentration is not a simple exponential function.

The rate of increase in local concentration as the peak concentration approaches becomes progressively less for stations further from the release point. This can be accounted for by the fact that as the peak concentration approaches a given station the size of the peak is decreasing. So long as the rate of approach of the peak concentration is faster than the rate of decay of the peak, the local concentration will increase with time. For stations well below the release point, the decrease in peak concentration and the increase in spread may combine to produce a rather long interval just before the passage of the peak concentration when the local concentration will remain nearly constant. In later tests some local concentrations reached their peaks even before the peak concentration of the contaminant distribution had passed.

The data are insufficient to show conclusively the form of the local decrease in concentration with time for the later tidal cycles; although other evidence indicates that the local decay curve is not exponential for large times.

Test 4 was run under the same conditions as Tests 1-A and 1, except for river flow, which was set at the relatively low value of 3000 sec ft at Trenton. The general features of the contaminant distribution curves, as shown in Figure 8, were similar to those found for Tests 1-A and 1. However, not only was the rate of downstream movement decreased due to the decrease in river flow, but the rate of decrease in peak concentration is appreciably less in Test 4 than for Test 1-A or Test 1. Figure 9 shows the change in peak concentration with time for Test 4. The rapid decrease in concentration during the first tidal cycle is similar to that which occurred in the earlier tests. The plot of the logarithm of concentration versus the time in tidal cycles after the second tidal cycle also

appears as a straight line, as in the case of Test 1-A and Test 1. However, the slope of this line in Test 4 is only one-half the slope of the decay curves for the earlier tests.

The movement of the peak concentration with time for Test 4 is shown in Figure 10. Again we see that the peak concentration moves at a slower rate than the calculated net water movement, though the differences are less here than in Tests 1-A and 1.

The local variation of contaminant concentration with time is shown in Figures 11 and 12 for selected stations above and below the release point. These figures also reveal the slower rate of decrease in the contaminant concentration for this test as compared to Tests 1-A and 1.

The rate of river inflow thus appears to exert a significant influence upon the rate of decrease in the contaminant concentration.

Tests 2 and 3 - Release Point 292.5

Station 292.5 is well into the estuary with salinities of about 15‰ for mean river flow. Test 2 was made with mean tide and mean river flow (12,350 sec ft at Trenton) while Test 3 was made with mean tide and high river flow (31,300 sec ft at Trenton).

Unfortunately, much of the data from Test 2 is unusable since large amounts of dye were adsorbed on the sample bottles. In addition, the density of the dye being less than the density of the water at the release point the contaminant was seen to "mushroom" and spread out in the surface layer during the earlier tidal periods. (In Test 3 the density of the dye was adjusted to that of the surrounding water and this difficulty was not

repeated.) Only the data for tidal cycles 6, 24, 36, and 42 were usable.

The longitudinal distributions of the contaminant concentration for various tidal cycles for Test 3 are shown in Figure 13. The distributions recoverable from the data for Test 2 were similar to those for Test 3.

Figure 14 shows the movement of the peak concentration for Test 2 as well as the computed mean downstream water movement. In Test 2 where the dye "mushroomed" into the surface layers, the downstream movement of the peak concentration was greater than or equal to the computed net water displacement during the first ten tidal cycles. This would be expected since the surface layers have an average downstream motion greater than the average for the entire body of water. As vertical mixing progressed, the motion of the peak concentration decreased.

Figure 15 shows the movement of the peak concentration and computed net water displacement for Test 3. Here again the movement of the peak concentration is slower than the calculated net water flow. The difference for this test was the greatest observed.

Figure 16 plots peak concentration against time for Test 3. It also shows the usable data from Test 2. Although the data from Test 2 show more scatter than the data from Test 3, the rate of decrease in peak concentration seems to be about the same for both tests. As in the previous tests the relation seems to occur in two parts, an initial rapid decrease followed by a slower exponential decay.

Since so much data were lost in Test 2 a conclusive evaluation of the effects of changed river flow on concentration is not possible. However, it seems that the effect at Station 292.5 may not be so great as at Station 52.5.

Curves showing the local change in contaminant with time in Test 3 at stations above and below the release point are given in Figures 17 and 18 respectively.

Tests 5, 6, and 7 - Release Point 111

These three tests were conducted to determine the effects of change in tidal amplitude on the time change in contaminant concentration. The dye volume, with a concentration of 1000 ppm, was introduced at high water slack at Station 111 which is located near Chester, Pennsylvania. This station is very nearly the upper limit of salt water intrusion for the conditions of mean river flow under which these tests were run.

Test 5 was conducted under conditions of mean tide, Test 6 under conditions of spring tide, and Test 7 under conditions of neap tide. Since the tidal velocities and tidal excursions would vary under these varying tidal regimes, it would be expected that the diffusion, and hence the time change in concentration, would vary among these three tests. As will be pointed out below the expected variation was neither clearly established nor definitely disproved by the data.

The longitudinal distributions of the contaminant at various tidal cycles for these three tests are given in Figures 19, 20, and 21. These distribution curves are similar to those encountered in previous tests in that they have the general shape of a normal function, and the change in peak concentration with time is large at first, becoming smaller for

later tidal cycles.

The downstream movement of the peak cor. entration was essentially the same for all three tests, and is shown in Figure 22. As with all the other tests, the observed movement of the peak is at a slower rate than the computed downstream water movement.

Figures 23, 24, and 25 are semilog plots of the peak contaminant concentration versus time for each of the three Tests 5, 6, and 7 respectively. Subsequent to the first two tidal cycles, the semilog plots of peak concentration versus time could be adequately described by a straight line for the first five tests. For Tests 5, 6, and 7, however, it appears that this simple exponential time variation is inadequate. In Figure 24 the curve from Test 1-A is also entered to show the sharp difference between the characteristic time change in concentration of the first five tests as compared to these last three tests.

The local time variation in concentration at selected stations is shown for these three tests in Figures 26, 27, 28, 29, 30, and 31.

The plots of peak concentration versus time do not show a systematic difference from test to test which can be readily ascribed to the differences in tidal regimes. One hypothesis concerning the effect of tide is that with higher tidal velocities mixing would be increased and hence the peak concentration should decrease at a faster rate, at least during the early stages. It is difficult to measure accurately the actual peak concentration at the first tidal cycle, and also after about 30 tidal cycles. Some information is obtained by comparing the peak concentrations for

the three tests for tidal cycles after the 1st and prior to the 30th. In Table I the peak concentrations for the 3rd, 6th, 12th, and 20th tidal cycles subsequent to release are given for each of the three tests.

Table I

Effects of Tidal Regime

Tidal Cycle	Concentration in ppm for Tests 5, 6, and 7		
	<u>Test 5</u> Mean Tide	<u>Test 6</u> Spring Tide	Test 7 Neap Tide
3	5 .4 5	4.58	5.81
6	3.25	2.83	3.34
12 -	2 . υ9	1.42	1.75
20	1.05	0.76	0.84

The concentrations for all four tidal cycles are lowest for spring tide conditions when mixing should be at a maximum. The concentrations for tidal cycles 3 and 6 are highest for the neap tide conditions, as would be expected, but for tidal cycles 12 and 20 the concentrations are highest for Test 5, which was run under mean tidal conditions. The tidal mechanism for the model failed after cycle 30 during Test 5, and it may be that a slight malfunctioning occurred prior to the final failure, resulting in somewhat erroneous results. In any case the data presented in Table I, while not conclusive, do suggest that the decrease in concentration is at a more rapid rate the greater the tidal velocities, at least during the early stages of mixing.

Summary Discussion

The composite information from all eight tests leads to the following general statements concerning the distribution and rate of change of
distribution of the dye contaminant.

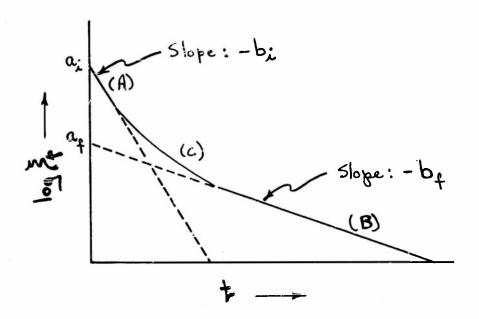
- 1. Within the first tidal cycle subsequent to release, the contaminant, which was in all cases introduced at high water slack as a cylindrical volume having a prototype diameter of 740 feet, spread over a longitudinal distance approximately equal to a tidal excursion (about 40,000 feet) and laterally from shore to shere.
- 2. After the first tidal cycle the longitudinal distribution of the contaminant approximated a normal curve.
- 3. Lateral and vertical variations in the contaminant concentration were observed during the early stages of each test. The sampling procedure did not allow a detailed investigation of the lateral or vertical variations in concentration, though the pertinent features that were noted are listed below:
- a. During the early stages of mixing, relatively concentrated segments of the contaminant volume were observed to become entrapped by eddies within small embayments or behind projecting marine structures. These entrapped contaminant segments would then feed out contaminant into the main stream after the main mixing volume had been carried out of the immediate area by the tidal currents. It appeared that this feature played a major role in contributing to the horizontal spread of the contaminant.

- b. At prominent bends in the river the advanced portion of the contaminant volume was observed to cling closer to that shore having a more rapid downstream movement of water.
- c. In Test 3, which was conducted in that part of the estuary which exhibits a net two-layered flow pattern similar to that described by Pritchard (1952) for the Chesapeake Bay, the upstream spread of the contaminant was observed to occur primarily along the bottom. During the early stages of the tests the vertical distribution for stations above the release point showed increasing concentration with depth, while those below the release point in general showed decreasing concentration with depth. After a few tidal cycles, vertical mixing had destroyed these general vertical variations in contaminant concentration.
- 4. The change in peak concentration is characterized by a very rapid rate of decrease during the first tidal cycle followed by a less rapid rate of decrease. In the first five tests, namely 1-A, 1, 2, 3, and 4, the variation in peak concentration subsequent to the first tidal cycle can be represented by a simple function of time. In Tests 5, 6, and 7, however, the variation of peak concentration subsequent to the first tidal cycle cannot be adequately represented by a simple exponential rate. The reason for the difference between these groups of tests is not at present evident.
- 5. The downstream movement of the peak concentration is significantly slower than the calculated velocity of the mean downstream water movement.

Theoretical Considerations

A Preliminary Analysis of the Characteristic Features of the Flushing Study in the Delaware Model

The semilog plots of peak concentration versus time in tidal cycles for Tests 1-A, 1, 2, 3, and 4 (Figures 4, 9, and 16) are all of the form shown in the following diagram:



Since this is a semilog plot the straight lines indicated by (A) and (B) may be expressed formally using the slope-intercept form of the straight line as

$$\log \sum_{i} = -b_{i}t + \log a_{i}$$
and
$$\log \sum_{i} = -b_{f}t + \log a_{f}$$
 respectively

where $-b_i$, $-b_f$ are slopes and log a_i , log a_f are the intercepts. Since

 $log A = B implies e^{B} = A$, we may write:

$$\sum_{\mathbf{p}, \mathbf{f}} = \mathbf{a_i} e^{-\mathbf{b_i} t}$$
 for the initial period
$$\sum_{\mathbf{p}, \mathbf{f}} = \mathbf{a_f} e^{-\mathbf{b_f} t}$$
 for the final period

If we sum these two equations

$$= a_i e^{-b_i t} + a_f e^{-b_f t}$$

we may use the result as an approximation to some (C) the contribution of (B) to (A) is negligible and conversely. Within the transition zone, (C), (A), and (B) will both contribute and the sharp corner will be rounded off.

Therefore, formally at least, peak concentration as a function of time may be written

As pointed out above, a peculiarity of this function is that for values of t outside of the neighborhood of t = 1 only one of the terms makes a major contribution to the value of . Thus during the first tidal cycle subsequent to release, the very rapid rate of decrease characterized by the first term of equation (1) predominates, and we have

(2)
$$= a_i e^{-b_i t}$$
 (first tidal cycle only)

After the second tidal cycle, this first term becomes unimportant, and the change in peak concentration is characterized by the final rate of decrease given by

(3)
$$= a_f e^{-b_f t}$$
 (after the second tidal cycle)

The variations in peak concentration for Tests 5, 6, and 7, subsequent to the second tidal cycle, do not appear to be adequately described by the exponential expression (3). For all tests, however, it appears that the expression

$$(4) \qquad \qquad \xi^{R} \frac{\partial \xi}{\partial t} = b'$$

holds. Here \underline{b} and \underline{n} are constants. The form of the solution of this differential equation depends on the value of \underline{n} . For the special case of $\underline{n} = -1$, the solution of (4) is

which, when b is identified with <u>-b</u>, is seen to be the expression which satisfied the observed concentration-time variations for Tests 1-A, 1, 2, 3, and 4.

For all \underline{n} other than n = -1, the formal solution of (4) is

(6)
$$b = bt + a$$
 , $b = b' (n+1)$

Designating (n + 1) by -m, this expression becomes

(7)
$$\sum_{p=0}^{m} bt + a$$

By solving for t, (7) can be expressed as

(8)
$$t = (1/b)$$
 $f^{-m} + C$; $C = -a/b$

or
$$\ln (t-c) = -m \ln \frac{c}{c}$$
 -lnb

Thus a plot of ln versus ln (t-c) should produce a straight line, of slope m. It is possible to obtain an approximate value of the constant c graphically from a plot of data which is suspected of showing a functional variation given by (8). This has been done for Tests 5, 6, and 7, and the resulting log-log plots of the concentration versus the sum of the time plus the constant c are given in Figures 32, 33, and 34. It is seen that the relationship is reasonably linear, and the equations for the three sets of data are:

Test 5 =
$$0.0341 \text{ t} + 0.225$$

Test 6 = $0.0513 \text{ t} + 0.0765$
Test 7 = $0.0506 \text{ t} + 0.114$

Since in equation (7) n = -m = 1, \underline{n} varies between -1.63 and -1.92 for Tests 5, 6, and 7, while for the five previous tests we found n = -1.

For the present we will confine ourselves to a discussion of the characteristic features of the decay expression

which can be expressed as

A plot of ln against time would result in a straight line of slope -b. The decay of radioactive material is represented by a function

of this type, and borrowing from the nomenclature of the nuclear field we may consider a characteristic time related to the rate of flushing of a contaminant which we will call the "flushing half-life." The flushing half-life is defined as the time interval required to reduce a given concentration to a concentration one-half as great. It should be noticed that if the rate of change of concentration is given by equation (9), then the half-life is a characteristic parameter independent of time or concentration; that is, if $\sum_{i=1}^{n} c_i c_i c_i c_i c_i$ represents the concentration at time $c_i c_i c_i c_i$, the same interval of time $c_i c_i c_i c_i$ is required to reduce the concentration to $c_i c_i c_i$, as is required to reduce the concentration from $c_i c_i c_i c_i$, to $c_i c_i c_i$, or from $c_i c_i c_i c_i$, to $c_i c_i c_i c_i$, to $c_i c_i c_i c_i$, to $c_i c_i c_i c_i c_i$, to $c_i c_i c_i c_i c_i$, or from $c_i c_i c_i c_i c_i$, to $c_i c_i c_i c_i c_i$, or $c_i c_i c_i c_i c_i c_i c_i$.

The half-life is readily obtained from the slope b, since

The flushing half-life is a convenient parameter for discussing certain features of the flushing tests. In Table II the half-life for the various tests and for the various characteristic portions of each test is given. It is seen that the half-life characteristic of the rate of concentration decrease during the first tidal cycle is only of the order of 1/100 of the half-life associated with the rate of concentration decrease subsequent to the second tidal cycle. Also we note that the half-life for Test 4, which was run under low river conditions, is two times the corresponding half-life for Tests 1-A and 1, which were run under con-

ditions of mean river flow, other conditions being the same for the three tests.

Table II

. 9	Half-life in tidal cycles		
Test	For first tidal cycle	After second tidal cycle	
1-A and 1	0.16	10.6	
2 and 3	0.11	13.5	
4	0.15	24.4	
5**	0.16	11.1	
6*	0.15	10.8	
7*	0.16	9.8	

^{*}Tests 5, 6, and 7 are included here for comparison, even though the peak concentration could not be adequately described by a simple exponential function of time. The calculations given are based on the concentration change from the fourth to the fortieth tidal cycles, neglecting the departure from the exponential form between these points.

Some Theoretical Considerations Suggested by the Model Study of Flushing

It can be shown (Pritchard, 1952, unpublished notes) that the onedimensional expression for the time rate of change of mean concentration in a channel of varying cross section may be expressed as

(13)
$$\frac{\partial}{\partial x} = -v \frac{\partial}{\partial x} + \frac{1}{A} \frac{\partial}{\partial x} \left\{ AK_{\xi} \frac{\partial x}{\partial x} \right\}$$

where x is the coordinate axis along the axis of the channel, v the mean

velocity in the x-direction, A the cross sectional area, and K the

aorizontal eddy diffusivity. If it is assumed that the contaminant dis-

tribution approximates the normal curve, then it can also be shown that

(14)
$$K_{\xi} = -\sigma^{2} \frac{\partial \ln \xi_{p}}{\partial t}$$

where \mathcal{F}_{p} is the peak concentration of the contaminant.

In Tests 1-A, 1, 2, 3, and 4 it was found that

Therefore, for these tests,

(15)
$$K_{\xi} = b \sigma^2$$

and the eddy diffusivity K varies as the square of the standard spread of the contaminant distribution.

In Tests 5, 6, and 7, however, it was found that

$$= a + bt$$
 equation (7)

for which the diffusivity would be given by

(16)
$$K_{\mathcal{S}} = \frac{\mathcal{B}}{A^m} O^{(2-m)} \text{ where } \mathcal{B} = \frac{b}{m} \left(\frac{M}{\sqrt{2^n}}\right)^m,$$

A is the mean cross sectional area of the segment of the estuary occupied by the contaminant volume, and M is the total weight of the contaminant.

From equation (16) it can be seen that, for these tests, the diffusivity varies inversely as the mth power of the cross sectional area and

directly as the (2-m)th power of the standard spread of the contaminant distribution.

In Tests 5, 6, and 7 m had an approximate range from 0.5 to 1.0 showing extremes for the diffusivity of

(17a)
$$K_{\frac{3}{2}} = \frac{\cancel{3}}{\cancel{A^0}} \cdot \cancel{5} \quad \boxed{1.5}$$
 and

Stommel (1949) has suggested that if <u>L</u> represents a scale factor characterizing the size of the area within which eddy diffusion is being investigated, then the eddy diffusivity is approximately proportional to the 4/3 power of <u>L</u>. This concept was developed for diffusion in an area unrestricted by lateral boundaries and would hence seem inapplicable to this study. It is interesting, therefore, to note that the mean exponent of for Tests 5, 6, and 7 is approximately equal to the 4/3 power suggested by Stommel.

That the movement of peak concentration was in all cases slower than the calculated net water movement calls for comment. Pritchard (1952, unpublished notes) has shown that, if the distribution of the contaminant approximates a normal curve, then the position of the peak concentration is given approximately by

(18)
$$m = D_{+} - \frac{M^{2}}{4\pi} \left\{ \frac{1}{A_{m}^{2} \xi_{p}^{2}} \frac{d \ln A_{m}}{dx} - \frac{1}{A_{o}^{2} \xi_{p}^{2}} \frac{d \ln A_{o}}{dx} \right\}$$

<u>m</u> is the position of the peak concentration when the origin x = 0 is set at the position of the peak concentration at time t = 0. D is the net

movement due to the velocity of the water, i.e. $D_{\underline{z}} = \int_{0}^{\infty} vdt$ where \underline{v} is the mean water velocity. The subscript "o" indicates the value of the parameter at time zero. The meanings of the other symbols are as previously explained.

In this study the factor in braces in the second term of the right-hand number of equation (18) was always positive and therefore $m < D_{\xi}$ is to be expected.

The development of a suitable theoretical description of other aspects of these flushing tests is in progress. However, in view of the unexplained difference between the characteristic rates of decay for the first five tests and for the last three tests, further experiments with the model would be desirable.

Suggestions Regarding Flushing in the Prototype

The question now arises as to the extent to which the results obtained from the model might be applied to the prototype. In this regard we may consider certain features of the model verification.

The model was constructed according to Froude scaling. The model velocity structure and salt distribution which would result from Froude scaling alone would not be comparable to the prototype, since the Reynolds scale would be far from satisfied. In order to increase the turbulence, the model was artificially roughened. The extent of this artificial roughening was empirically determined by roughening until the velocity distribution and the tidal heights matched those known to exist in the prototype.

It was then found that the salinity distribution in the model satisfactorily matched that found in the prototype under various river flow regimes.

This salinity verification implies that the artificial roughening of the model successfully scaled those eddies in the turbulent regime which are important in maintaining the salt distributions. There is strong indication, however, that the scale of the turbulent eddy system which is involved in any particular mixing process depends on the size of the contaminant volume being mixed. Therefore, the eddies which would be involved in the mixing of a highly concentrated contaminant spread over only a small area may not be properly scaled in the model, even though the eddies important in maintaining the salt distribution are properly scaled from the prototype to the model.

On the basis of the above reasoning, the following preliminary conclusions regarding the application of the results of the model studies on flushing to prototype conditions are presented.

1. It is doubtful that the extremely rapid rate of concentration decreases observed during the first tidal cycle in the model studies actually represent in magnitude the corresponding initial rate of mixing which would occur in the prototype. Immediately after introduction of the dye, large gradients exist and molecular diffusion would be of some importance in the model, though less so in the prototype since there can be no scaling of this phenomenon from the prototype to the model. Likewise, as discussed above, the eddy scale important to the mixing of the

restricted contaminant volume during this early phase may not be properly scaled in the model.

Undoubtedly, the rate of concentration decrease during the early phase of mixing in the prototype will be substantially greater than at later stages, since the initial contaminant volume may spread laterally as well as longitudinally (which is the case in the model also). However, the rapid decrease in concentration in the model during the first tidal cycle is substantially greater than would be expected as a result of the coupling of the lateral spread with the longitudinal mixing. This is a feature which may allow prototype verification, since it is possible that tracer material which could be followed for one or two tidal cycles might be safely added to the natural estuary.

2. The rate of concentration change observed in the model subsequent to the second tidal cycle is probably applicable to the prototype.

By this time the contaminant had spread from shore to shore laterally
and over approximately a tidal excursion longitudinally. The eddy regime
contributing to the mixing would be of about the same order of magnitude
as that which is important to the maintenance of the salt distribution, and
hence probably properly scaled from prototype to model.

The flushing half-lives given in Table II for the period subsequent to the second tidal cycle after dye introduction are therefore probably representative of the expected rates of flushing in the prototype for contaminant distributions which are spread over approximately a tidal excursion.

3. The suggestion that the eddy diffusivity varies with at least the

first power, and possibly the second power, of the standard spread of the contaminant distribution throws some doubt on the application of techniques for computing flushing characteristics of an estuary which are based on the salinity distribution, or conversely, on the observed or computed distribution of fresh water. Though these techniques may adequately describe the fresh water-salt water distribution, they may not describe the concentration change of a contaminant which is introduced into the estuary with an initial distribution considerably different from the existing fresh water or salt water distribution.

Acknowledgements

Throughout my association with this project I have been impressed by the efficient and courteous assistance rendered by the U. S. Army Engineers Waterways Experiment Station. Particular mention should be made of the help given by Mr. Henry Simmons, Mr. William Robb, and their staff on the Delaware Model Project.

The U. S. Navy Hydrographic Office supplied the services of Mr. John Recknagel for all tests except the trial run. Mr. Recknagel handled the field project for the last four tests. His interest and conscientious work were of considerable importance to the successful completion of the project.

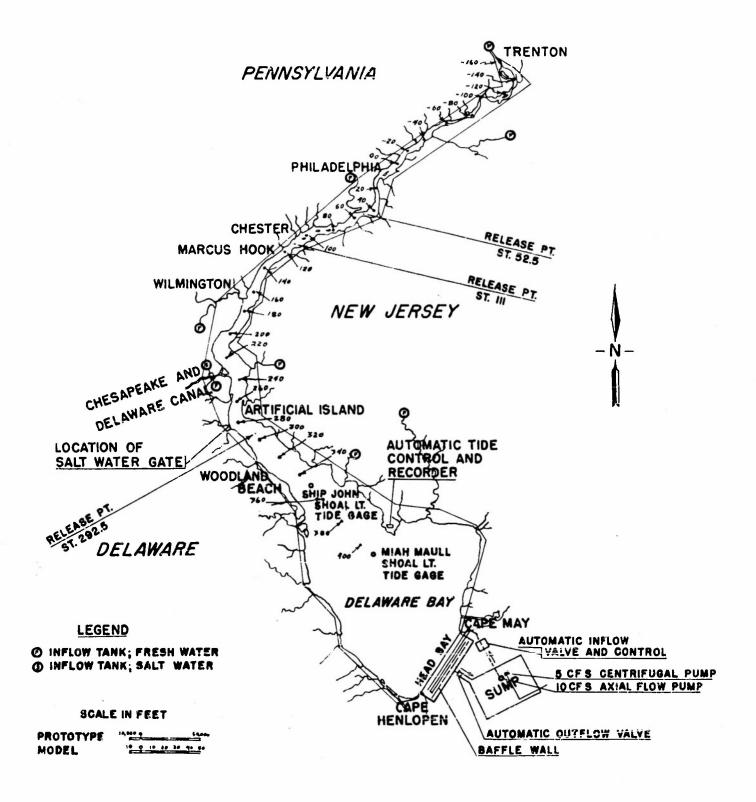


Figure 1 Location Map for Model Study of the Delaware Estuary.

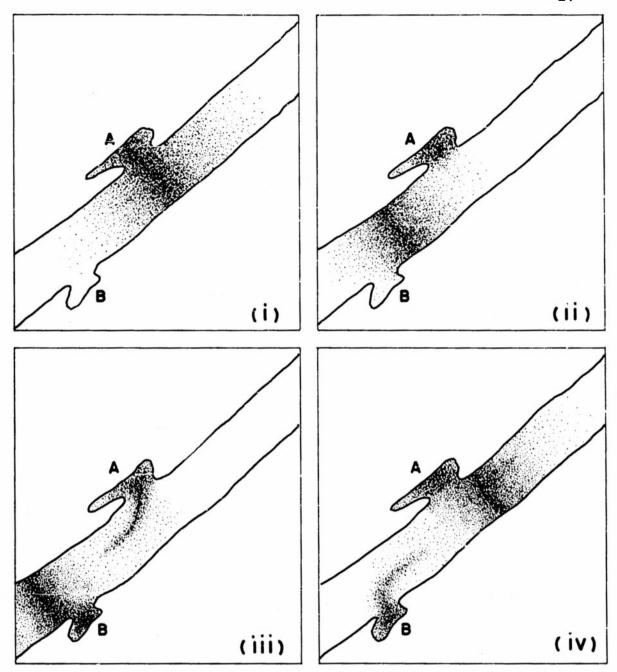
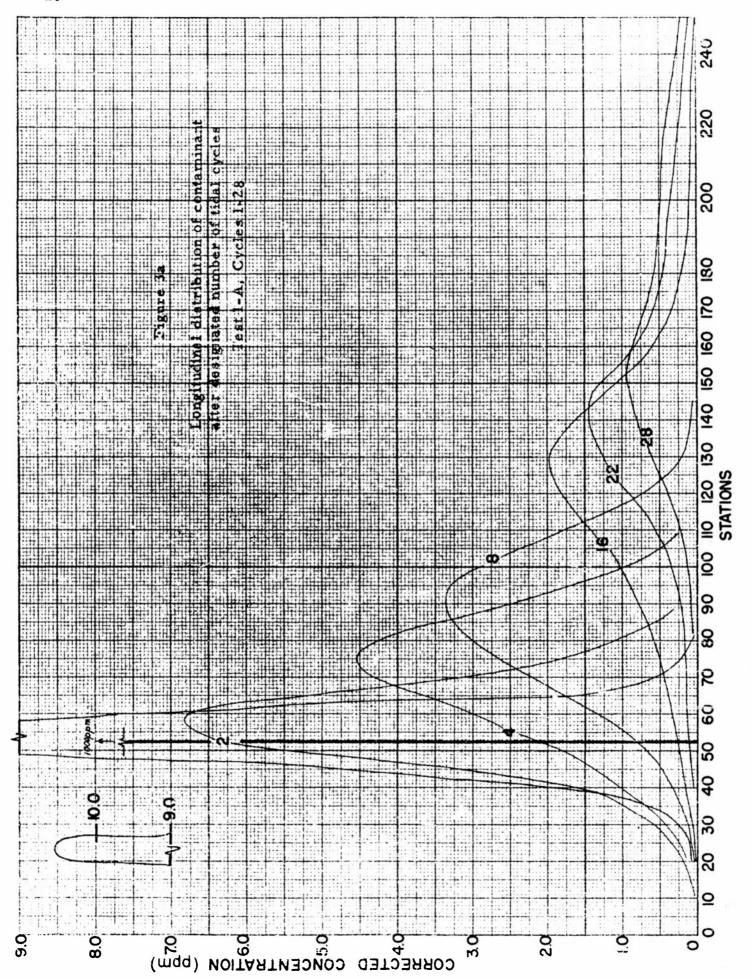
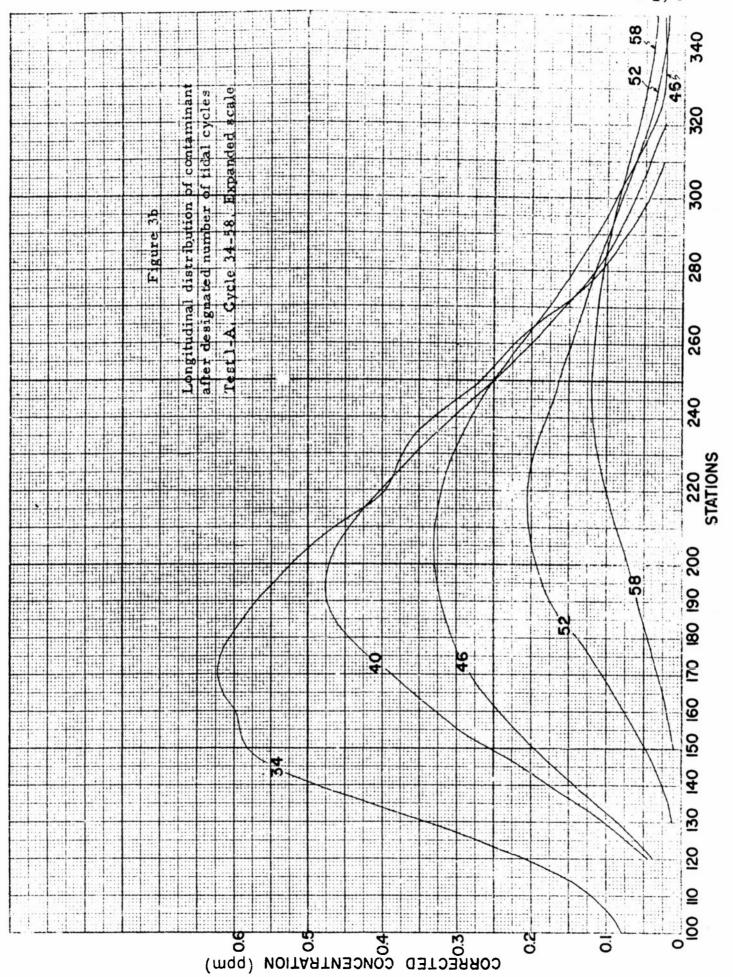
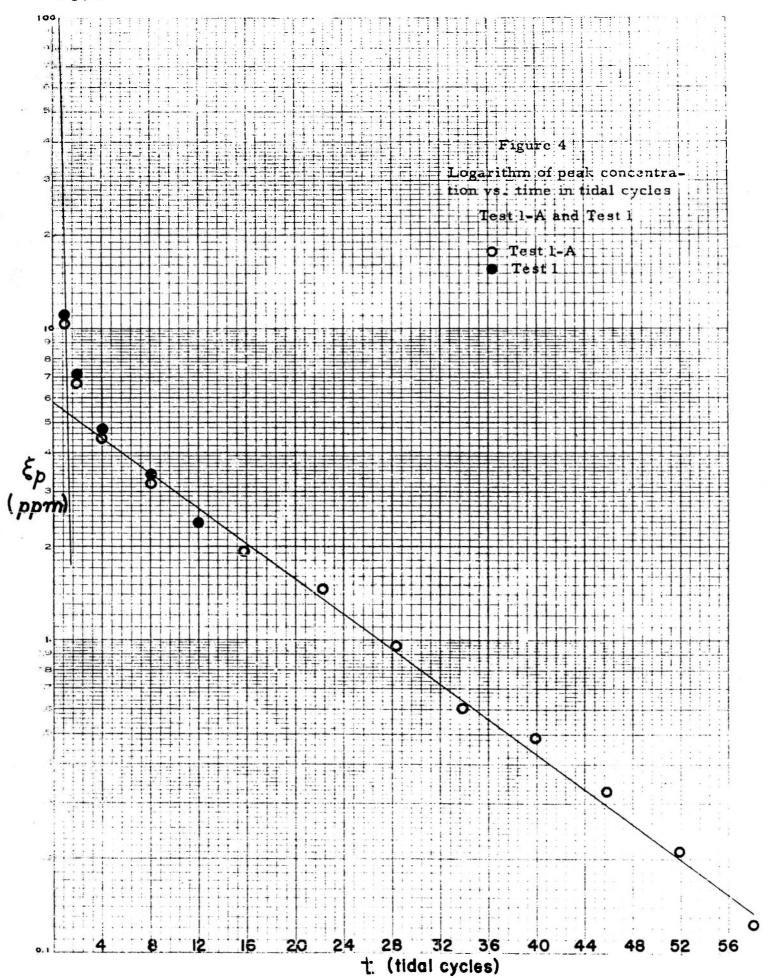
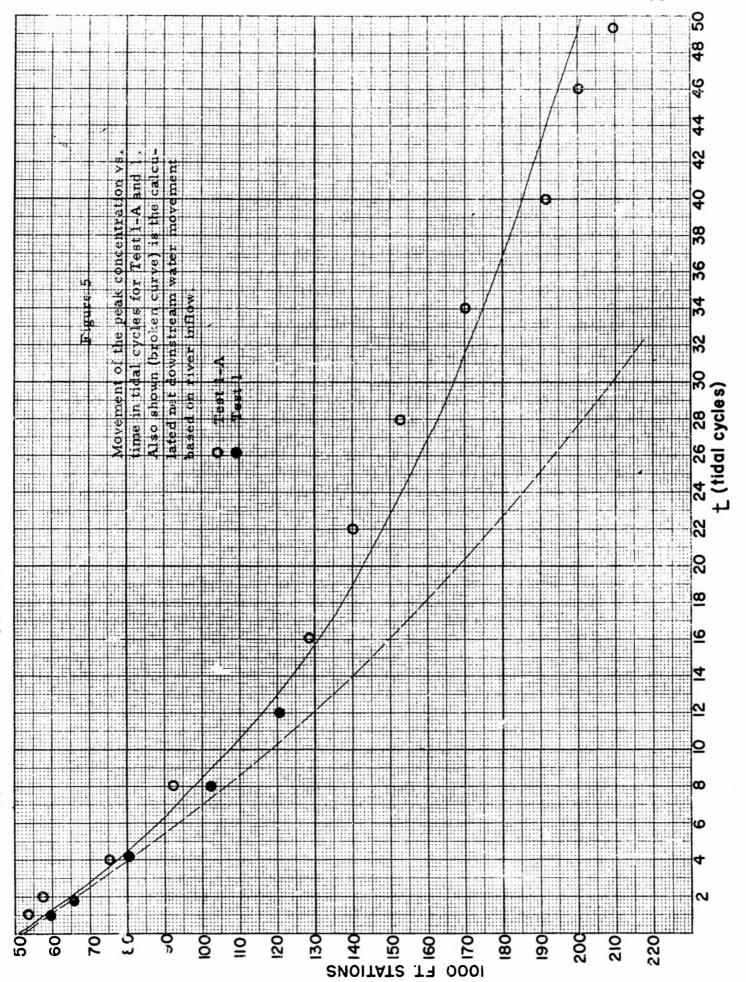


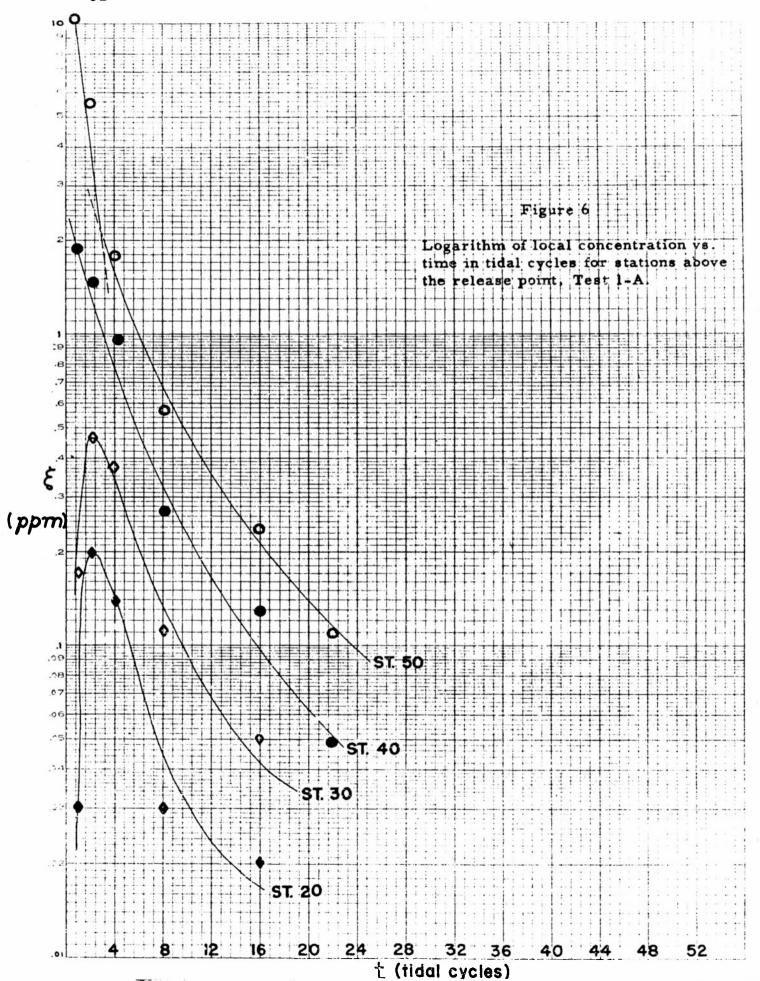
Figure 2. Schematic representation of dye entrappment by shore features. Concentration is indicated by intensity of shading. As peak concentration of main dye volume moves downstream on the ebb tide, a portion of the dye spreads into shore indenture A, as in diagram (i). This portion of the dye is entrapped by the indenture as the main dye volume moves on downstream, as shown in diagram (ii). In diagram (iii) the entrapped dye feeds out into the main channel, this contributing to the longitudinal spread. The main dye volume is shown in this diagram to have spread into a second shore indenture. In diagram (iv) the phenomena is shown repeated for a second indenture B on the flooding tide.

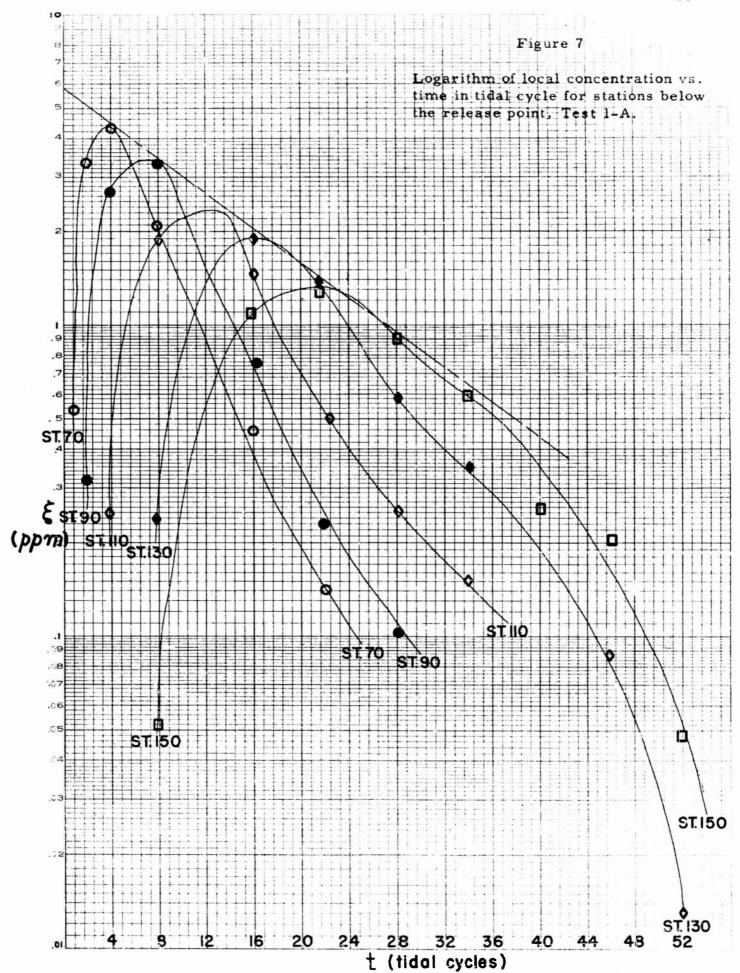


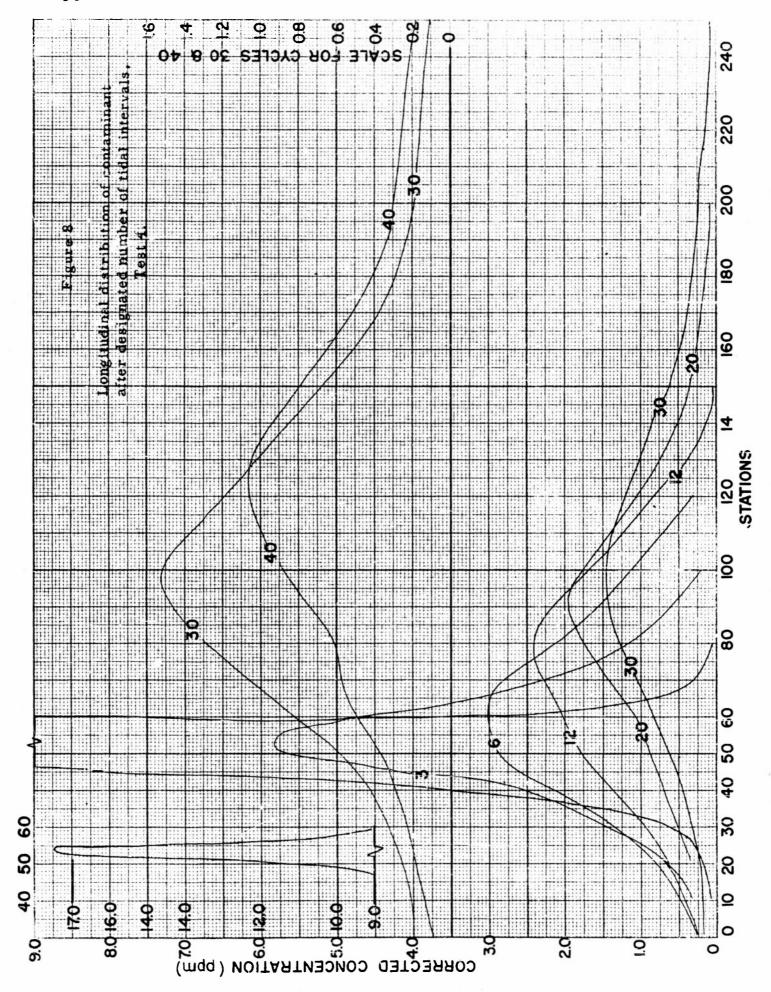


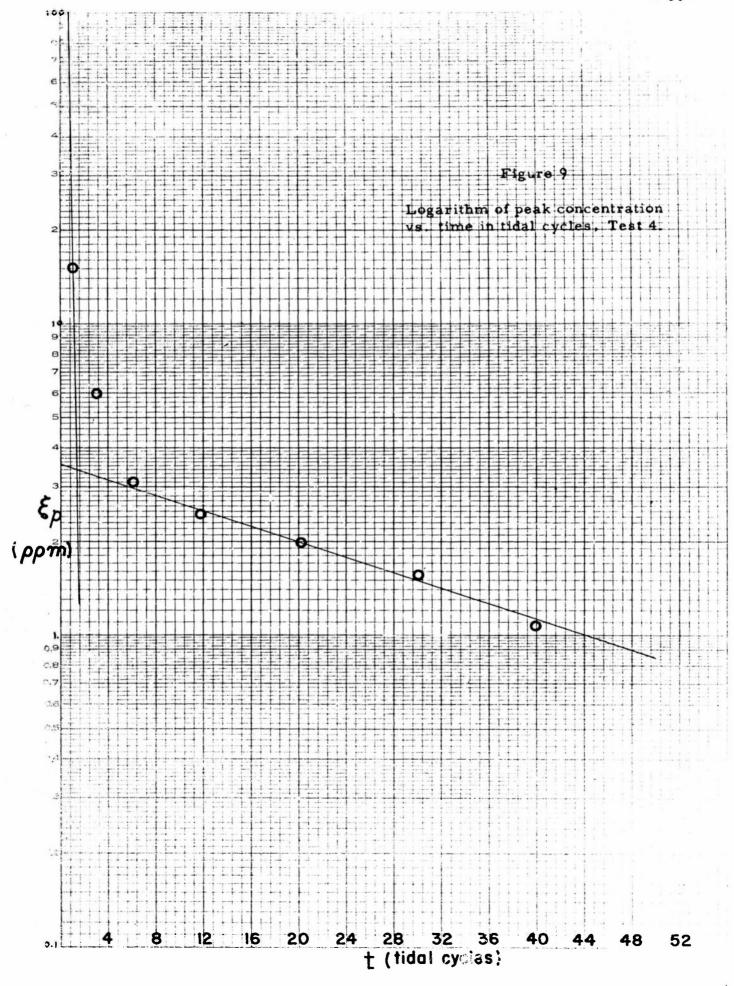


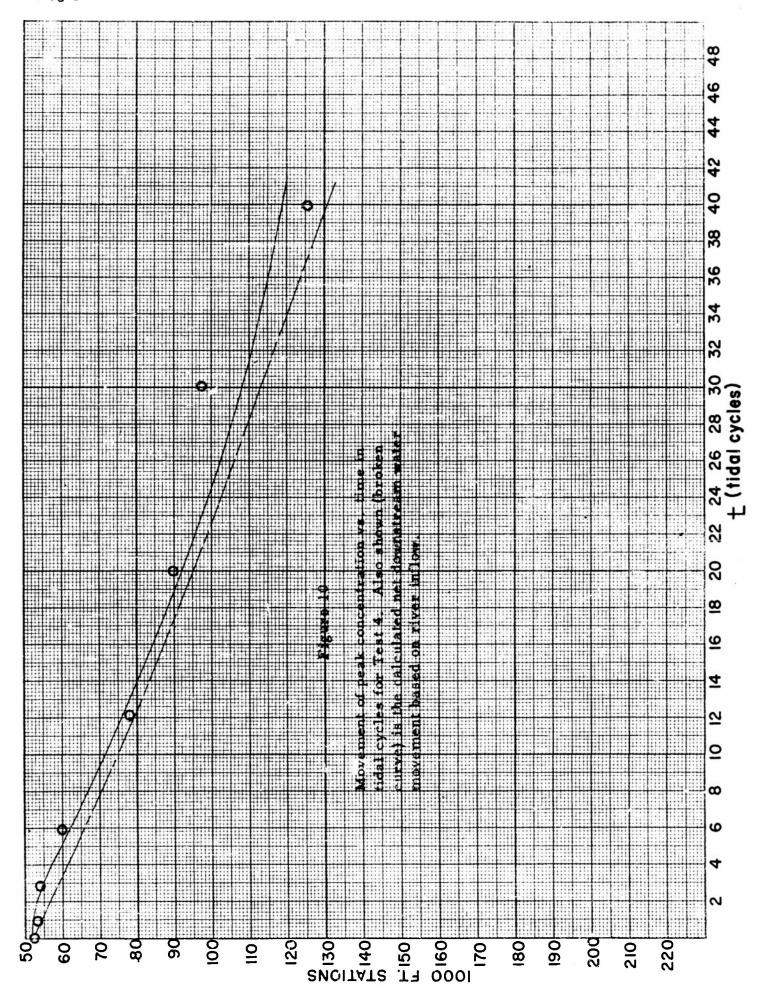


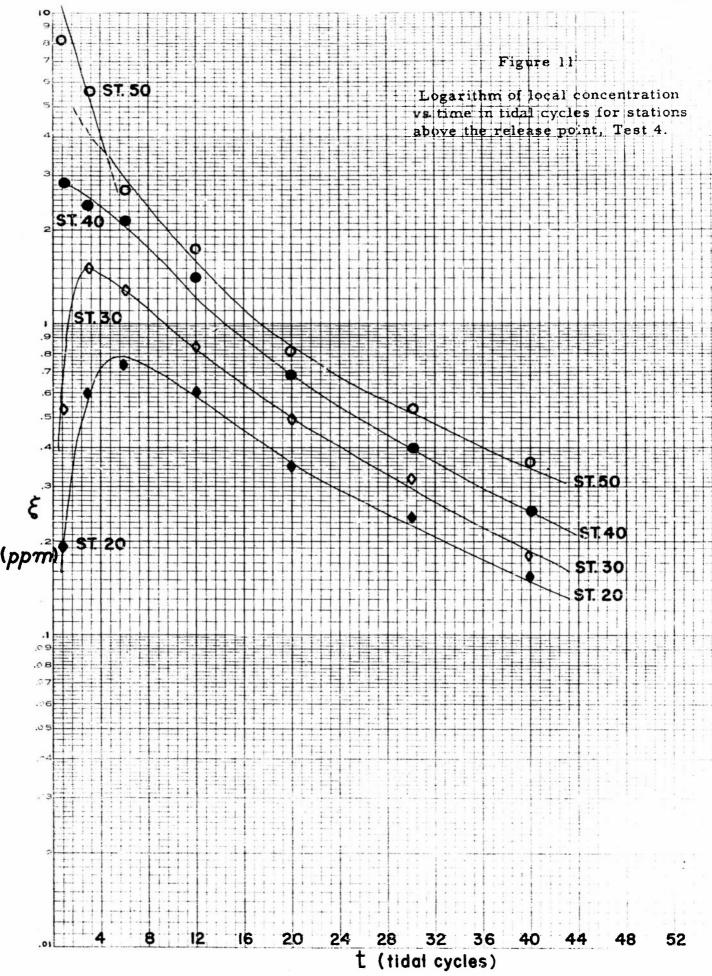


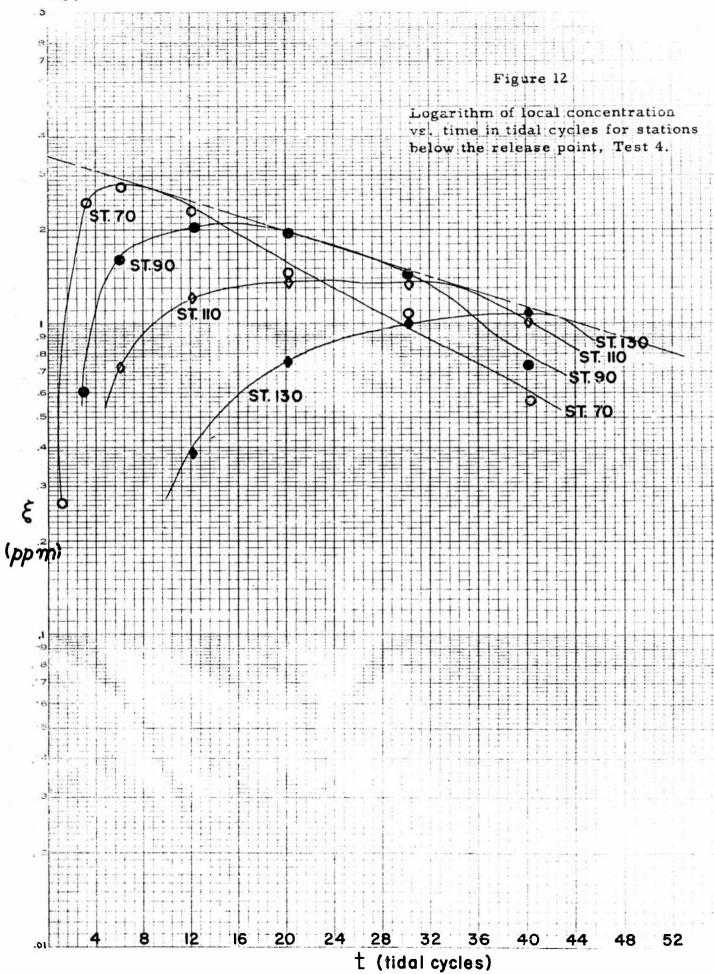


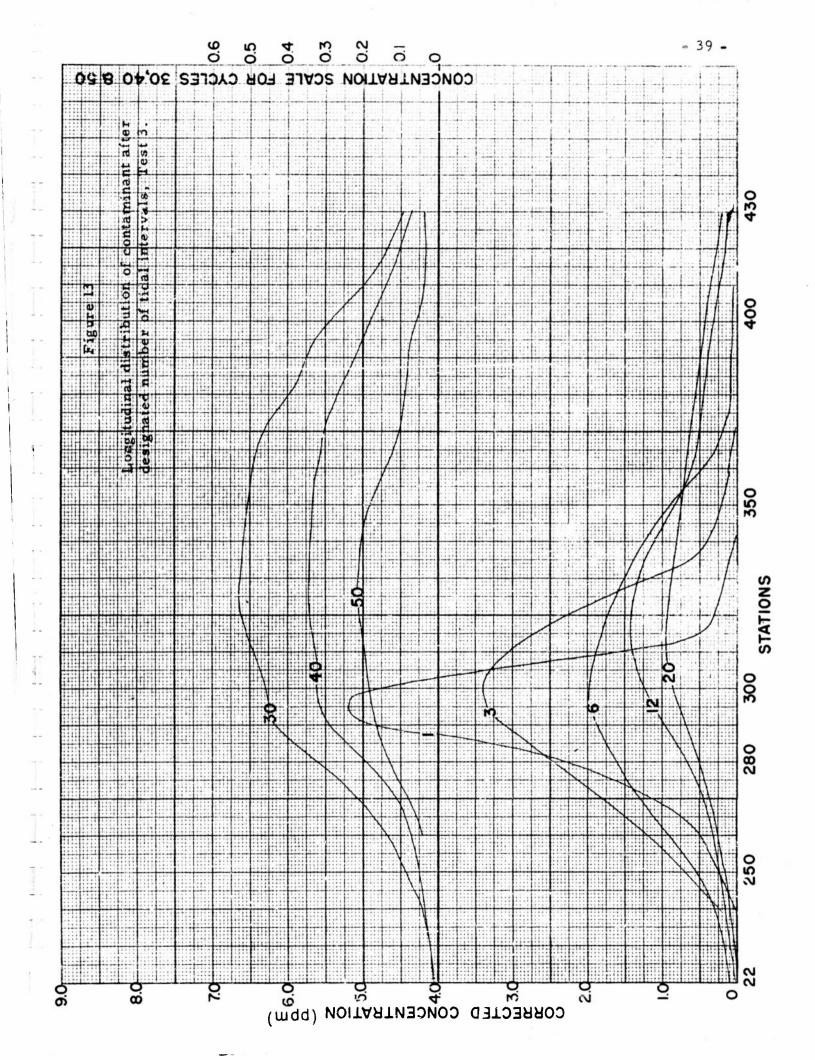


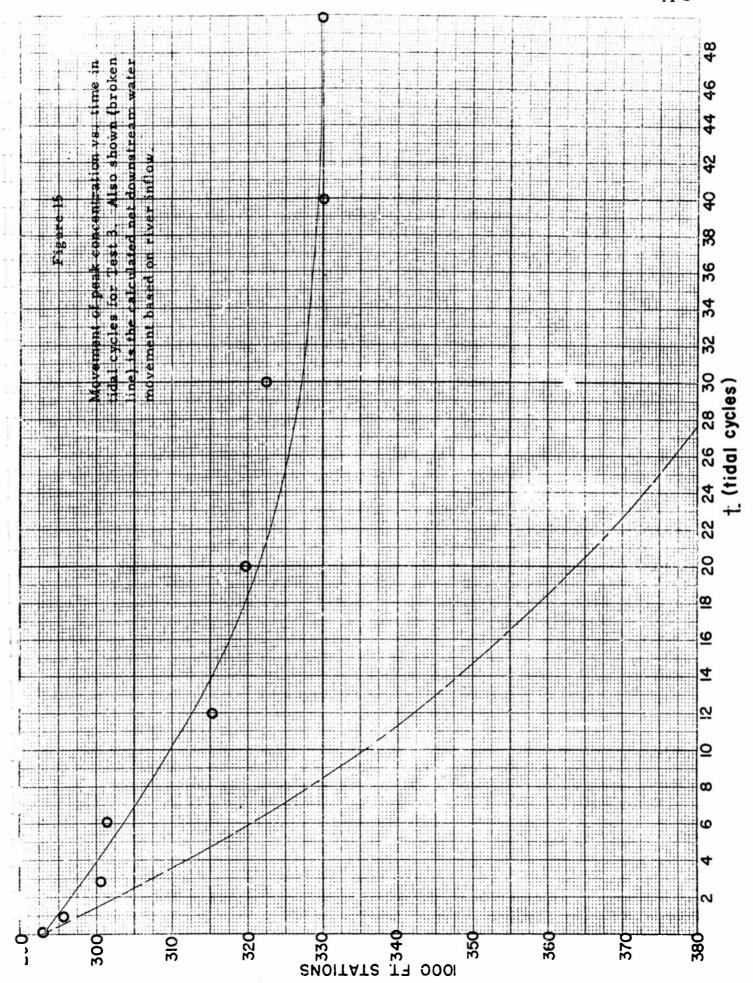


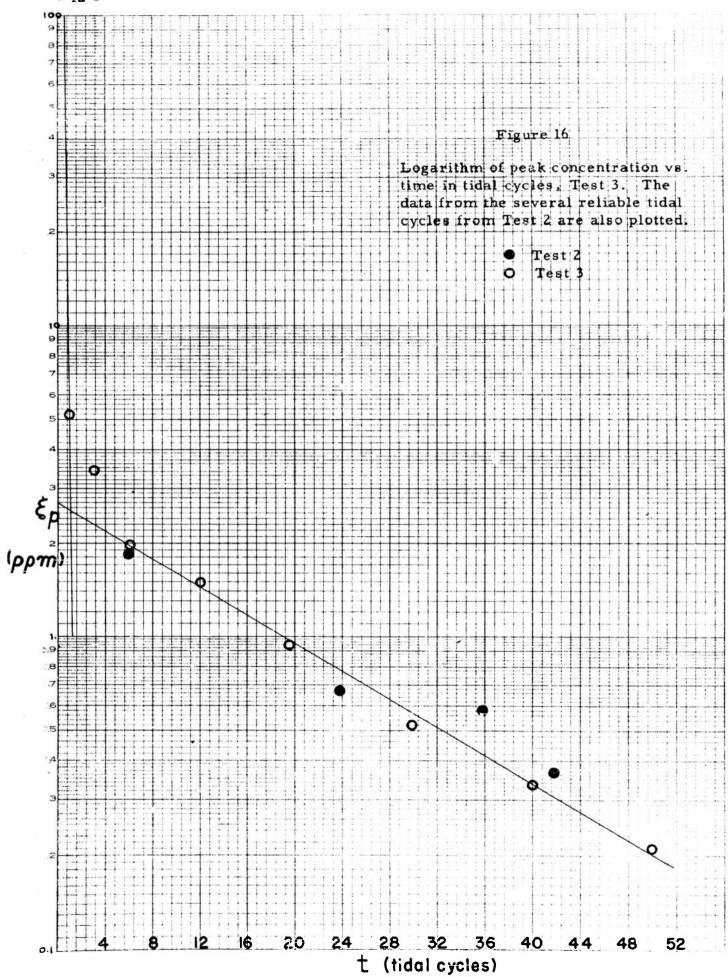












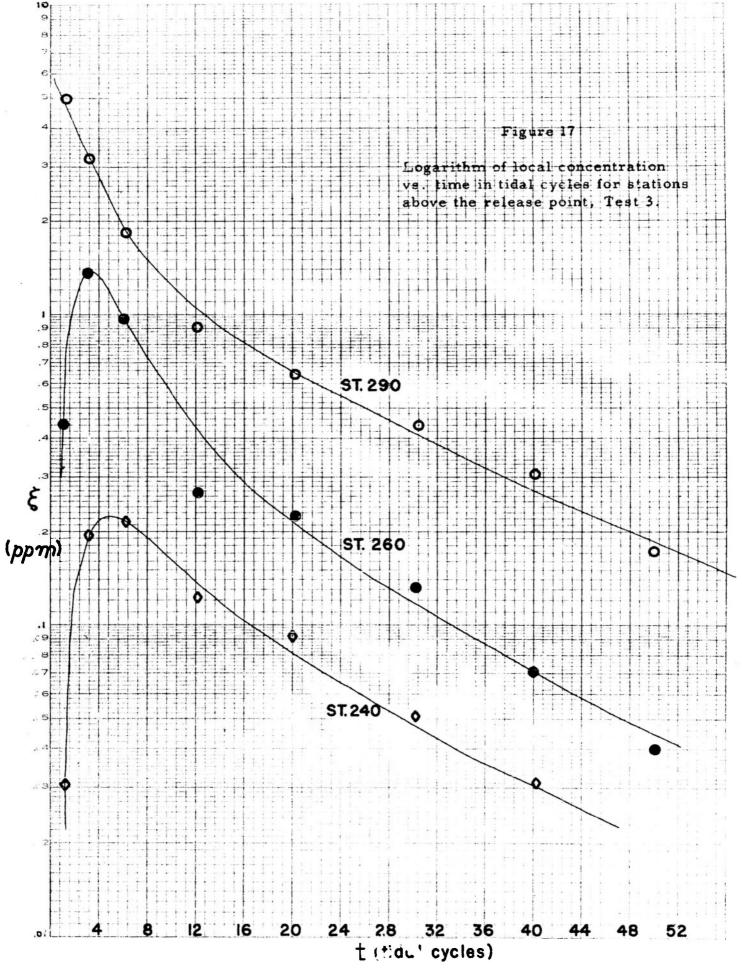
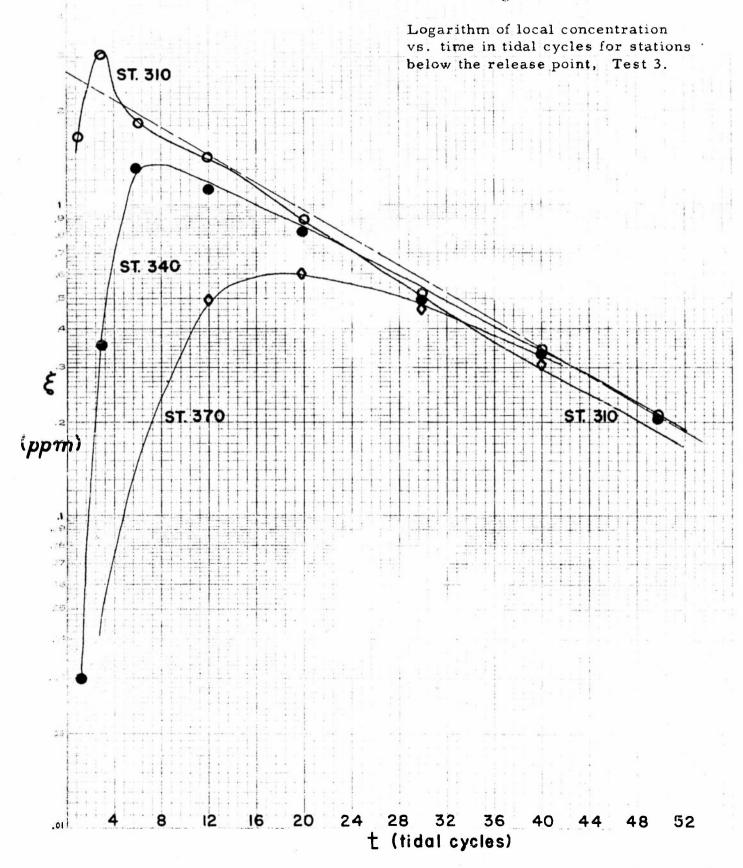
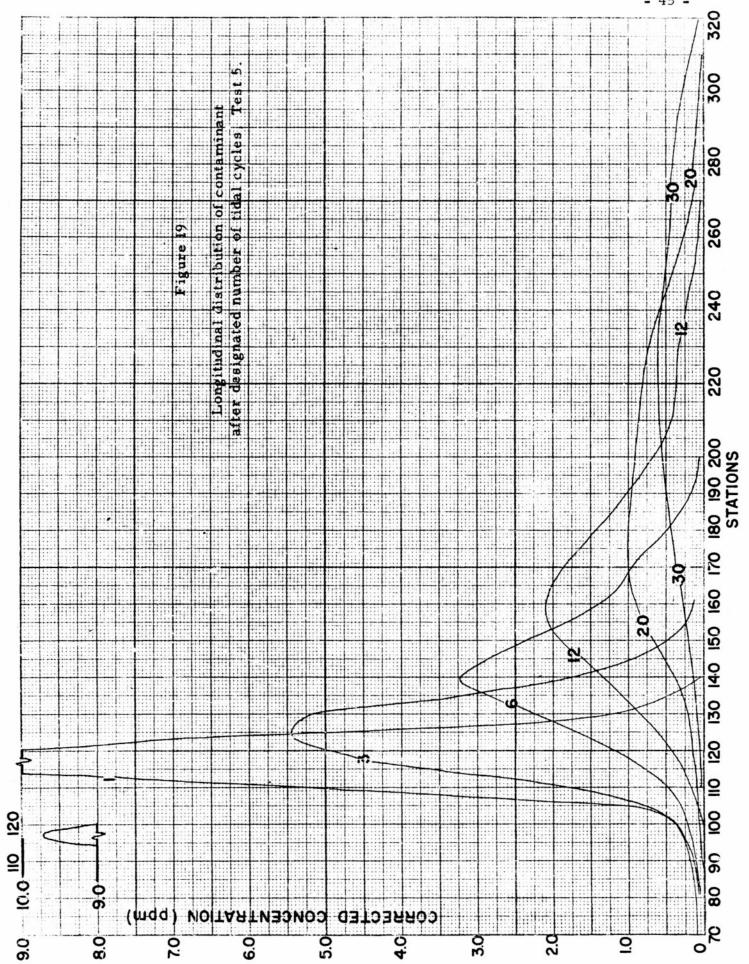
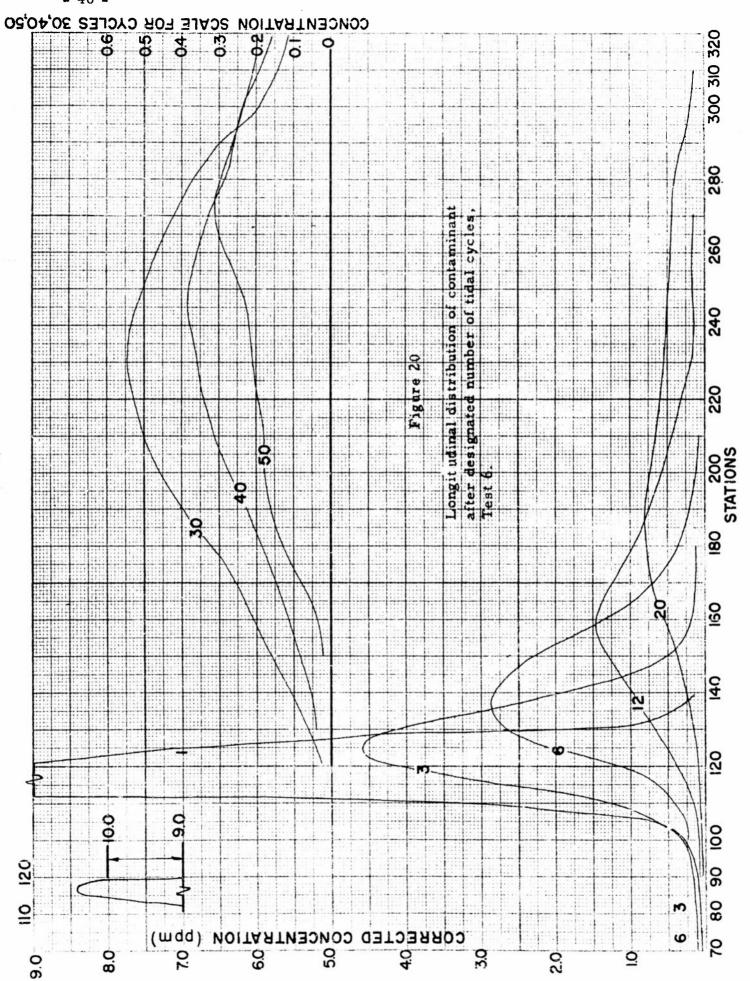
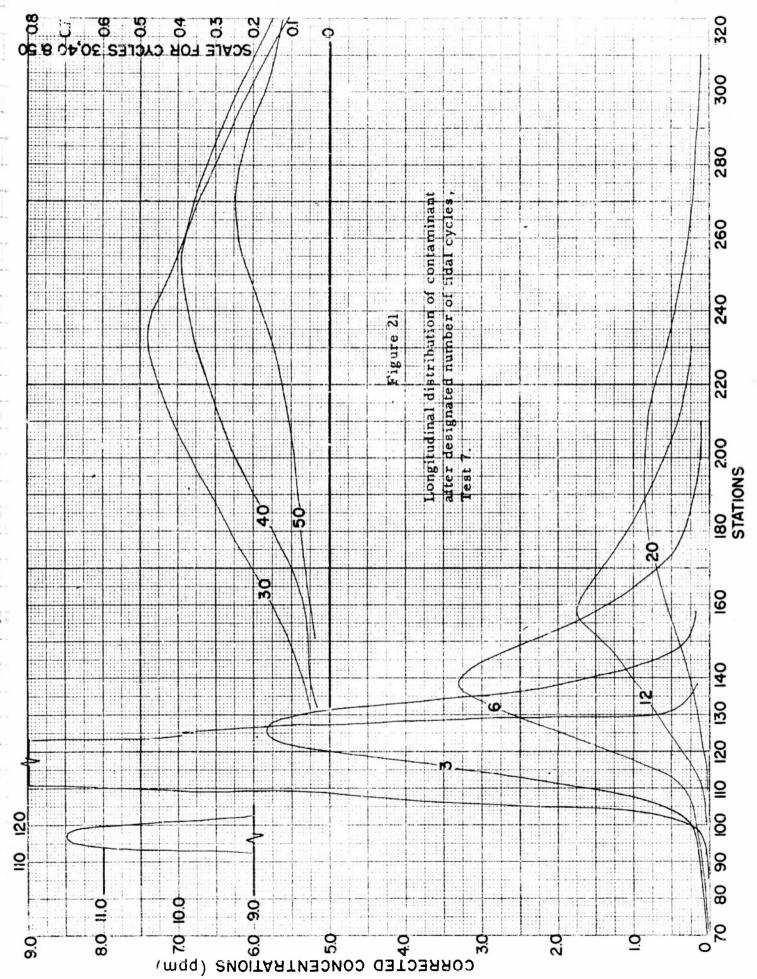


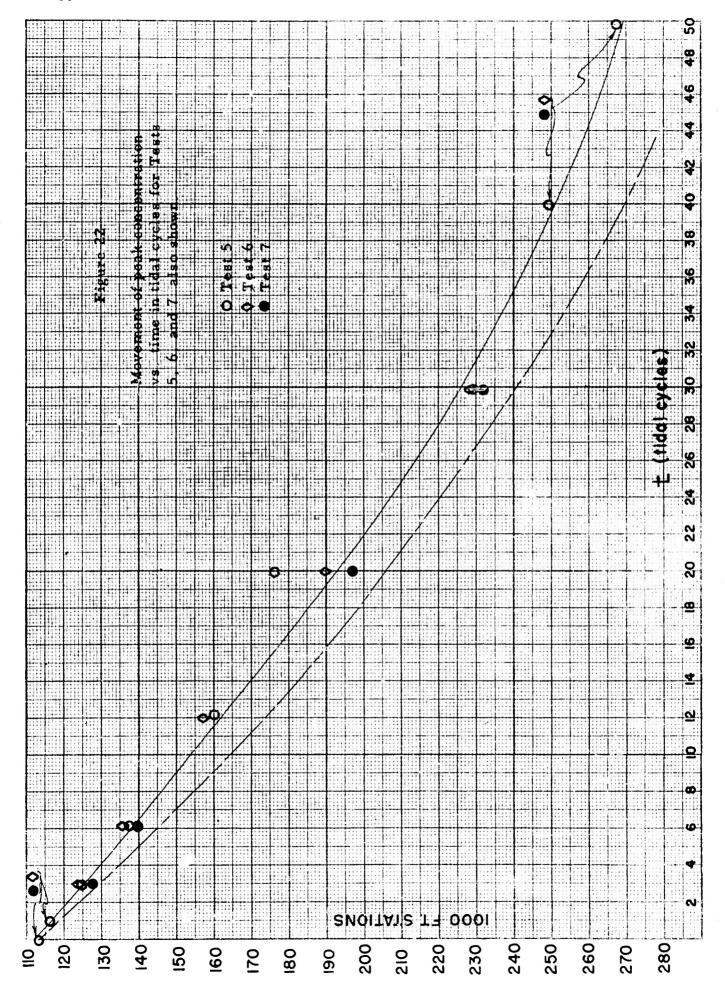
Figure 18

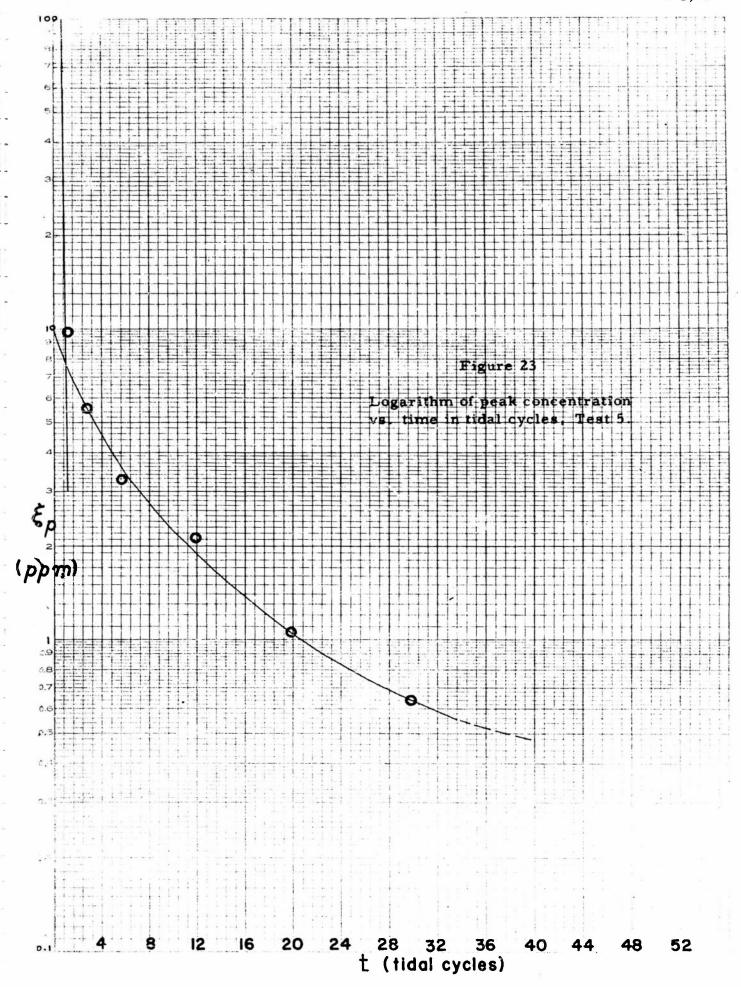


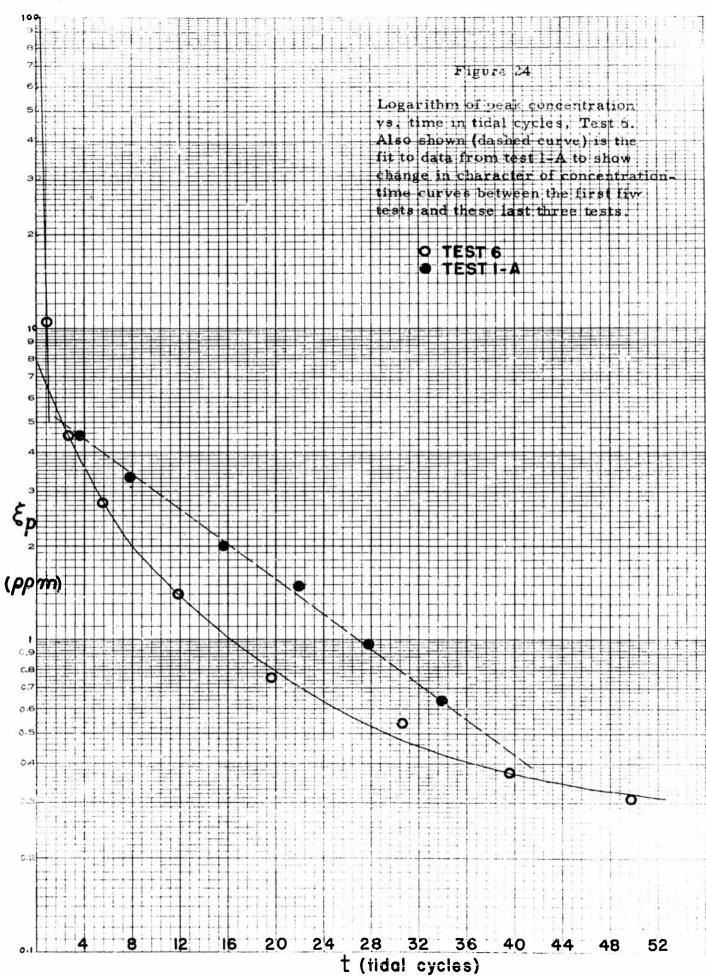


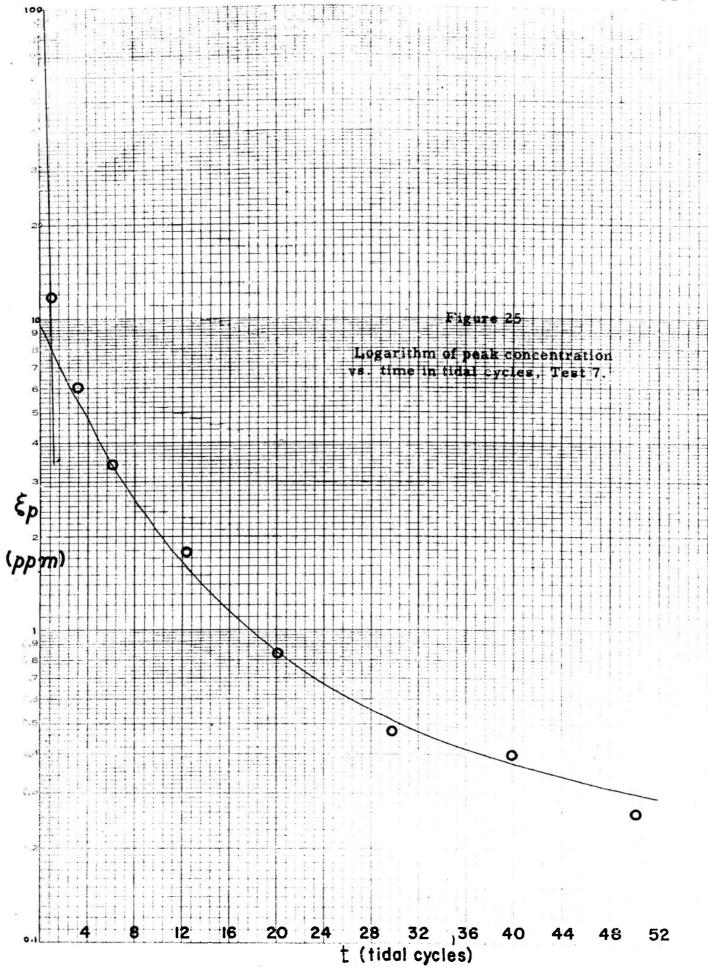


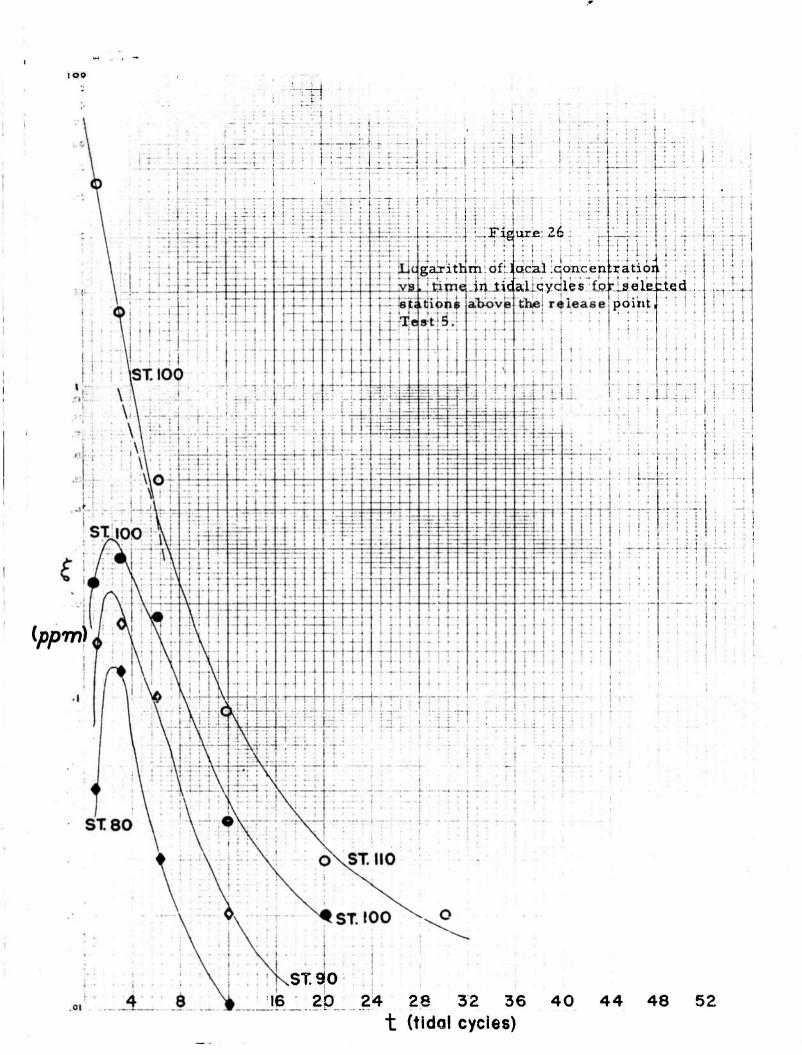


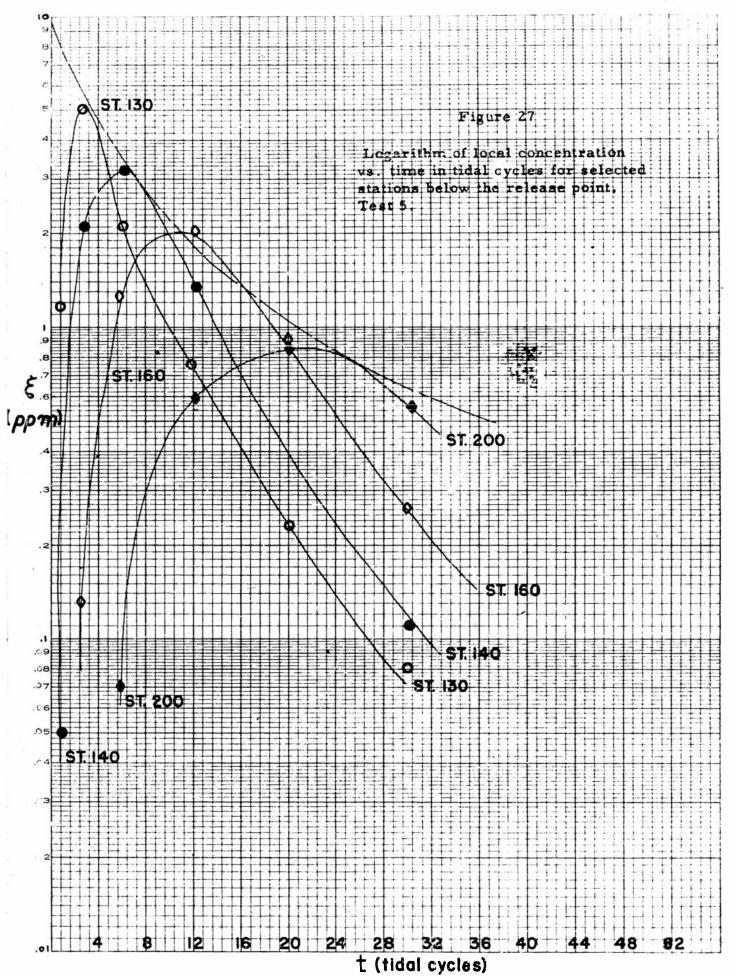


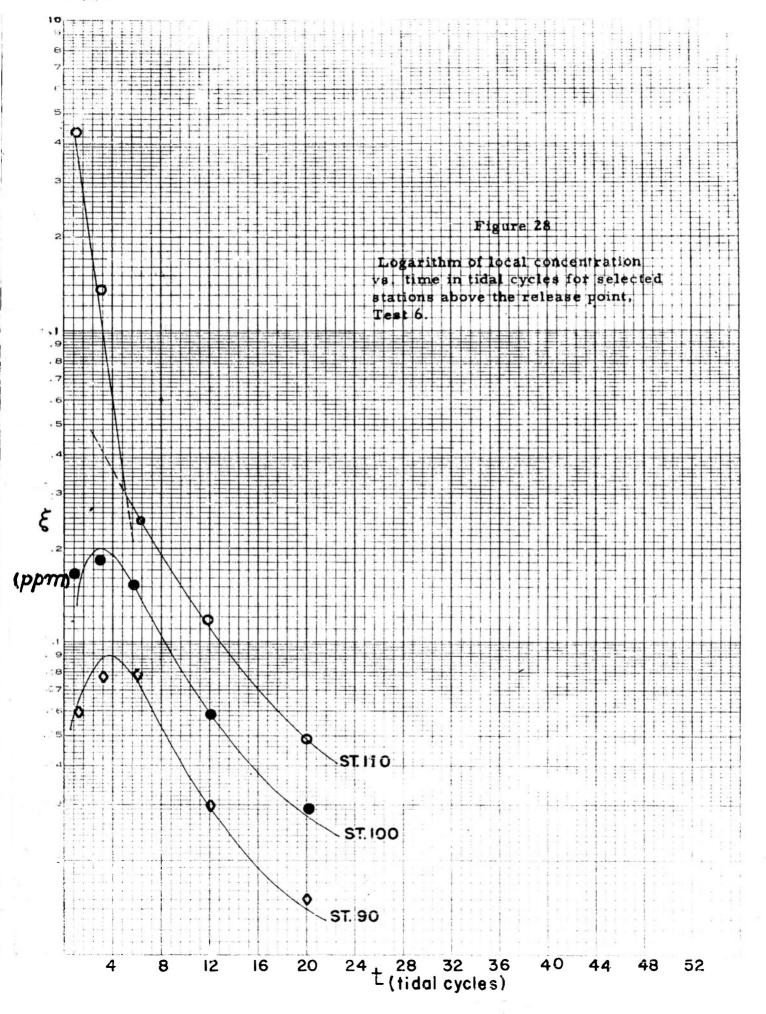


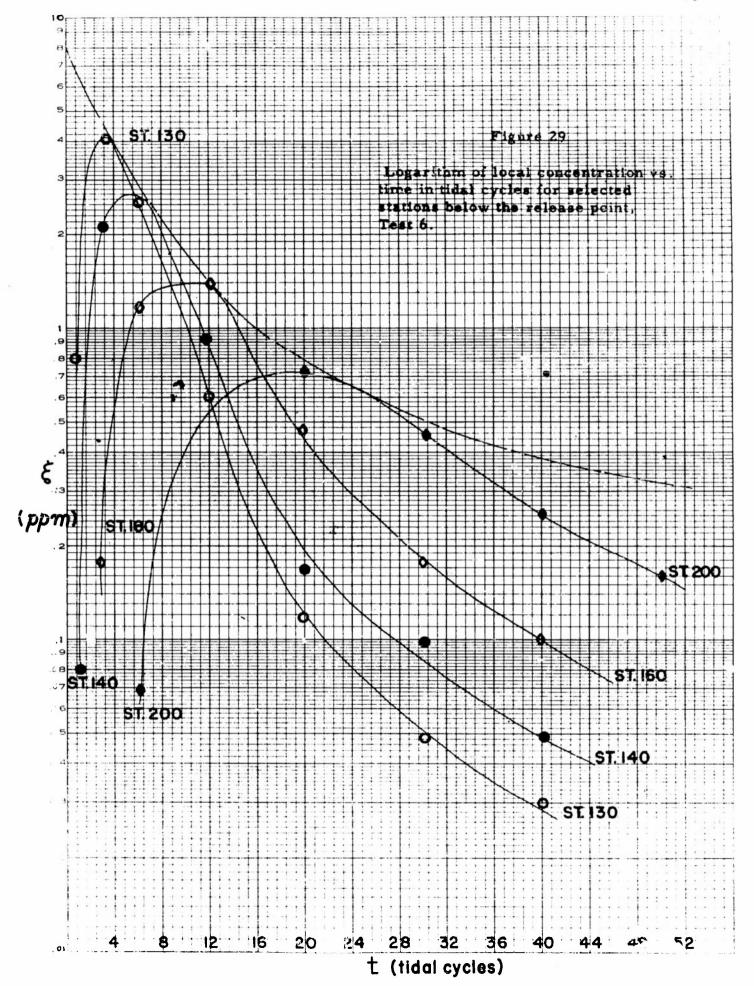


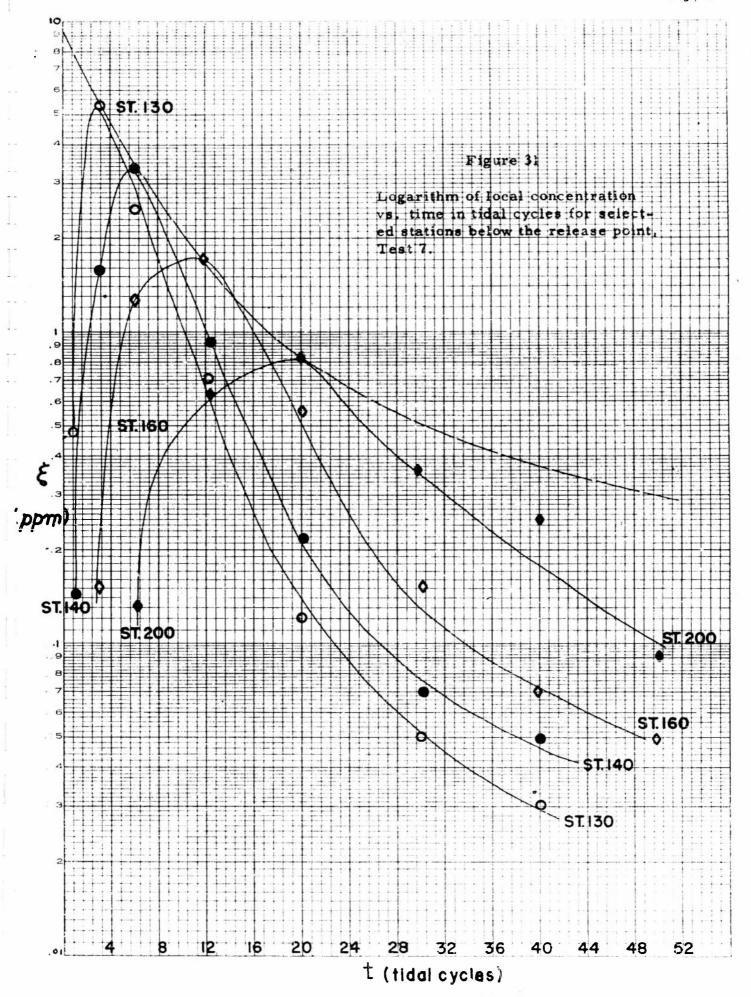


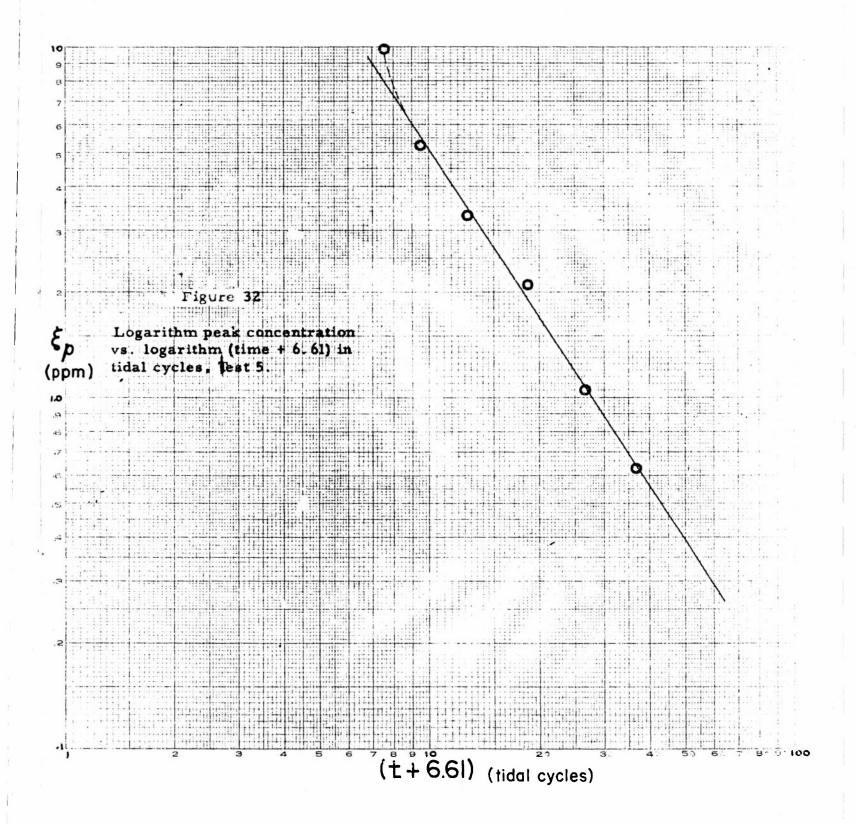


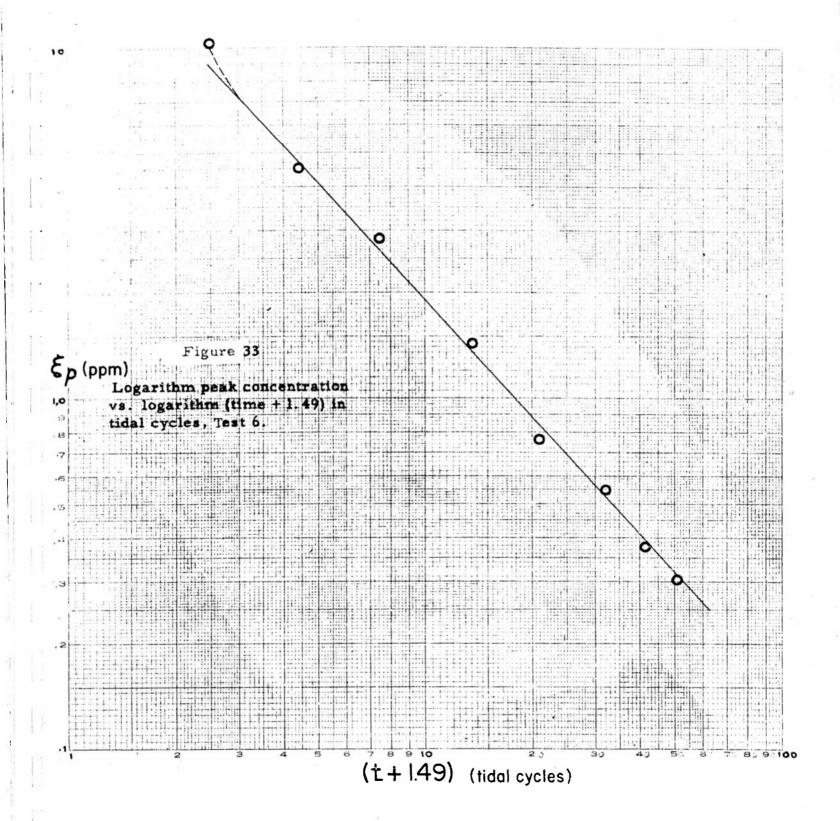


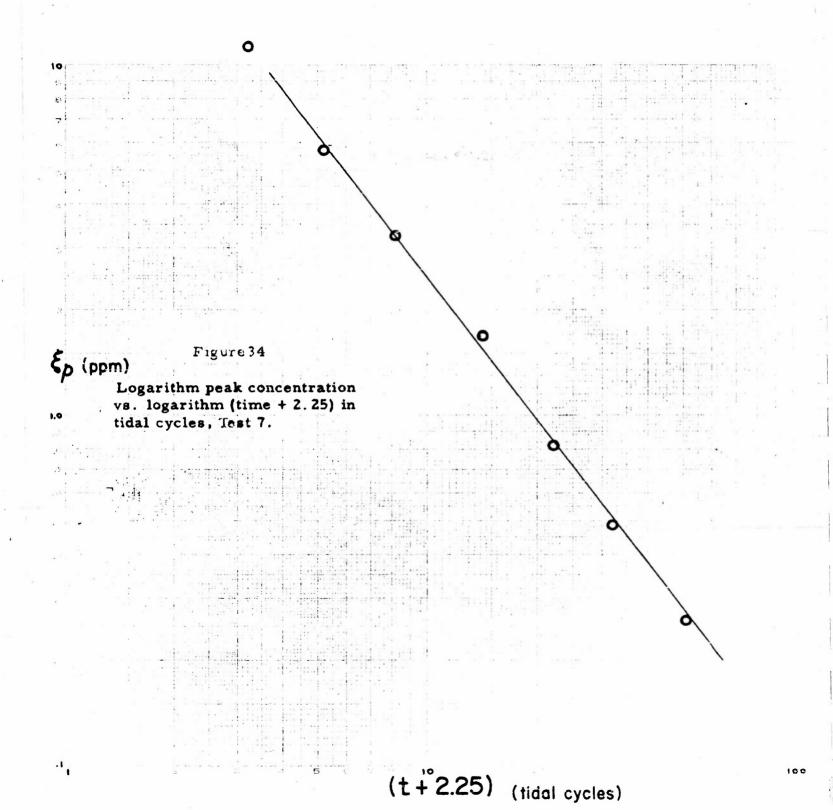












APPENDIX I

Description of the Dye Release Gear, the Water Sampling Equipment, the Method of Sampling, and the Determination of Dye Concentration

Dye Release Gear

The gear used to release the methylene blue used in these experiments is shown in Figure A-I-1. It consists of an open brass cylinder held firmly against a rubber lined base by means of a two inch wide bar and two outrigged rods. The rods were screwed into threaded holes in projections of the bottom piece, and pulled down on the two inch bar which was placed across the top of the cylinder. The bottom of the cylinder was thus drawn tightly down on the rubber sheet which was bonded to the base and leakage from the container was prevented.

At the appropriate release point a recess was made in the bottom of the model of just the correct size and shape to hold the bottom piece of the container. The depth of the water at the release point was accurately measured at high water, and the height of dye in the container was made to agree with this depth of water.

During the flooding tide just prior to the release time the dye container was introduced into the model, the bottom piece resting in the recession. The rods were unscrewed from the bottom and withdrawn, while one man held his foot securely on the rod across the top of the container, bearing down with sufficient force to prevent leakage of dye from the bottom of the container. At precisely high water slack of the tidal period chosen for release, the cylinder, now no longer secured to

the bottom piece, was lifted quickly but carefully upward from the model, leaving the bottom piece remaining in the recess in the model bottom, and the cylindrical dye volume suspended in the water of the model. Since the recess had been made to just fit the bottom piece of the container, the bottom of the model with the bottom piece in place was without any indentation or protrusion resulting from the release of the dye.

Sampling Equipment

Since the model has a vertical scale of 1 to 100, 0.01 foot of depth in the model corresponds to 1 foot of depth in the prototype. The sampling equipment, therefore, had to be made relatively small so as not to interfere with the flow.

The basic unit of the sampling gear was a thin glass tube, bent at a right angle near the end and drawn to an elongated point. The glass tubing was connected by plastic tubing to a 50 ml syringe. The assembly could be operated by hand as a single unit, or grouped together with other similar assemblies to give a multiple depth sampler. In any case the glass tubing was taped to a metal rod so that the metal rod projected at least a slight amount below the lowest glass tip.

In the single unit hand samplers, the glass tube was taped to the metal rod so that the bent tip was about 0.2 ft from the bottom of the rod corresponding to a prototype distance above the bottom of 20 ft. In this manner the rod could be placed with its end on the bottom at the desired sampling position in mid channel and the sample would be drawn

from about middepth, since the channel up to Philadelphia is maintained at the prototype depth of 40 feet.

The multi-depth sampling gear is shown in Figure A-I-2. The glass tubes were taped to a metal rod so that the bent tips occurred at appropriate intervals -- usually surface, middepth, and bottom. The rod was part of a tripod assembly usually employed in measuring the surface elevation of the water in the model, and the sampling assembly was thus supported in position by the tripod. The plastic tubes leading from the three glass tubes were connected to three 50 ml syringes secured to a wooden rack. The pistons for the syringes were all clamped to a wooden bar, which in turn was connected to a cable leading over a pulley at the top of the wooden rack. This cable was attached to a bar secured to three household rattraps.

A spring latch held the three rattraps, which were connected in the open or "set" position. In this position the cable was just taut with the pistons in the syringe all the way in. When the latch was released, the spring on the rattraps would pull down on the cable, and, hence, up on the syringes and the sample would be drawn.

In Test 1-A, use was made of a motor driven multi-sampler which took eleven simultaneous samples, drawing the sample evenly over the 7-1/2 minute tidal period of the model. This mechanism was not employed after the first test, since there was some difficulty in fitting the results from these samples with the samples taken at high water slack.

Sampling Procedure

Before introduction of the dye, the rattrap samplers were set up at appropriate stations above and below the release point. Other locations were chosen where single hand-drawn samples were to be obtained. On selected high water slack periods, subsequent to the release, samples were collected at the designated positions.

Since the tide in the Delaware is partially progressive, a particular phase of the tide occurs downstream first and can be followed up the Bay and the River. Small strips of paper were strewn on the surface of the model, and the time of high water slack determined at each particular station by noting he time when these floats stopped moving upstream. Thus, the sampling was not simultaneous at all stations, but related to the tidal stage, the first samples being taken at the downstream edge of the spread of the dye, and the last samples for any particular tidal cycle at the upstream end of the spread. With several individuals participating in the sampling, it was quite possible to keep pace with the progressing tidal phase as it moved upstream.

The samples were transferred from the syringes into wide mouth quart jars. As will be pointed out in a later appendix, this was far from the most satisfactory procedure. The jars were then assembled for analysis.

Determination of the Dye Concentration

The concentration of dye was determined photometrically using both a Beckman model DU spectrophotometer and a Coleman spectrophotometer.

At the beginning of each run a sample was taken from the dye volume to be introduced, and appropriate volumetric dilutions were made from this sample. The Beers-law calibration curve of adsorption versus concentration was then obtained for each instrument, using the diluted samples.

Samples of water from the model were also drawn prior to the dye introduction to serve as blanks in the calibration.

The ground around the model is quite dusty, and even though the model is well washed down between tests, there occurs in the model water some suspended material which appeared to interfere with the determination of low concentrations. The procedure employed in eliminating this difficulty is essentially the procedure described by Carritt and Carpenter (1951) in their paper on corrections for turbidity interference in chlorophyll analysis.

The procedure depends upon the contrast between the extinction curve due to the dye and the extinction curve resulting from the turbidity. This difference is shown in Figure A-I-3. The broken line A represents schematically the extinction curve for the dye in distilled water. This curve is characterized by a sharp maximum in the extinction at a wave length of 665 mm. The dashed line B is representative of the typical extinction curve resulting from turbidity. This curve shows a steady decrease with wave length, with no characteristic adsorption peak.

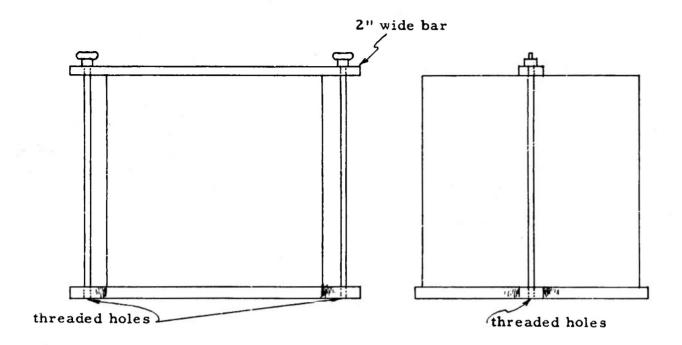
The addition of these two curves gives curve <u>C</u>, drawn as a continuous line. The typical extinction curve for the dye is moved up and modified somewhat by the turbidity.

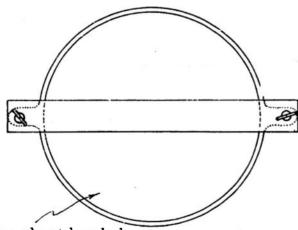
Normally the extinction E_m at the characteristic wave length n is taken as proportional to the concentration. However, if the sample is somewhat turbid, the resulting extinction E_m would indicate a larger dye concentration than actually existed. If two points, E_a and E_b are taken on the dye extinction curve well away from the peak extinction at wave lengths n and n respectively, and n is the linear interpolation between n and n at wave length n then the distance n to n is also proportional to the concentration. However, the distance n is approximately equal to the distance n on the combined extinction curve for coth dye and turbidity.

The procedure calls for running a calibration curve of the difference, $E_m-E_c \text{ versus concentration.} \quad \text{To do this, measurements are made of the extinction readings at the three wave lengths } \lambda_a$, λ_m , and λ_b . From the extinctions E_a and E_b the correction E_c can be obtained since

$$E_c = E_a - (E_a - E_b) - \frac{\lambda_m}{\lambda_b} - \frac{\lambda_a}{\lambda_a}$$

The difference $E_m - E_c$ is then taken as proportional to the concentration. The unknown samples are analyzed in the same manner, and hence the influence of the turbidity is essentially eliminated.

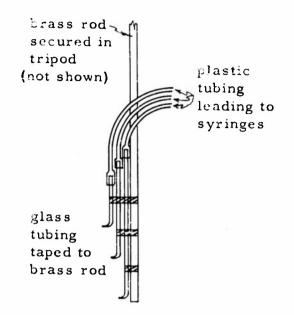




rubber sheet bonded to bottom piece

Figure A-I-l

Schematic drawing of dye release gear



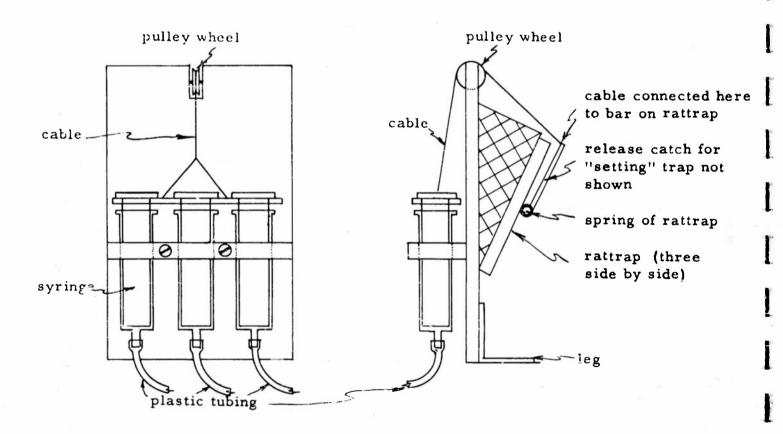


Figure A-I-2. Three depth rattrap sampler

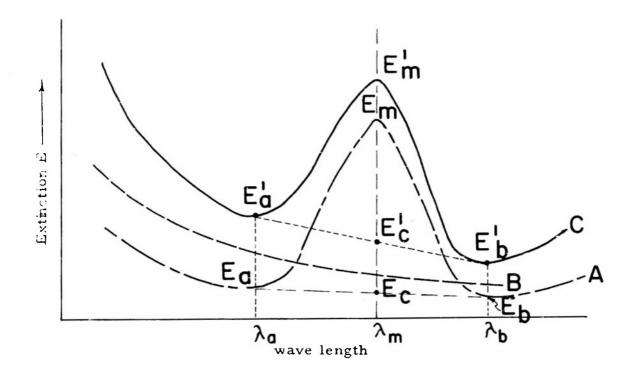


Figure A-I-3. Effect of turbidity (curve B) on the extinction of dye. The curve for the dye in distilled water is \underline{A} , and in water of turbidity given by curve \underline{B} the combined extinction curve is given by \underline{C} . The distance ($\underline{E}_m - \underline{E}_c$) is, however, approximately equal to ($\underline{E}_m' - \underline{E}_c'$), and may also be expressed as a function of dye concentration.

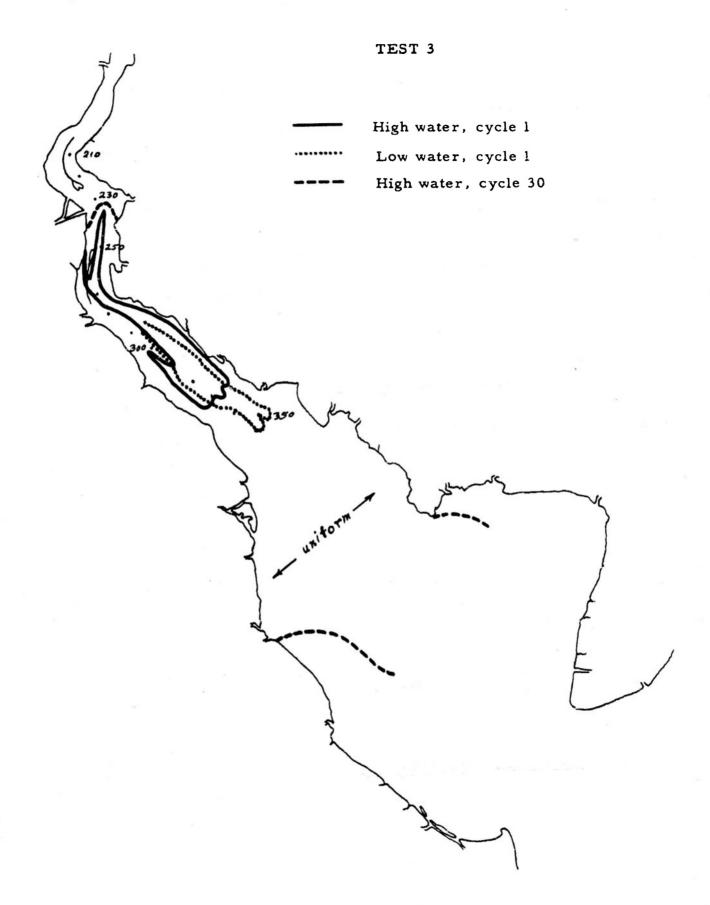
APPENDIX II

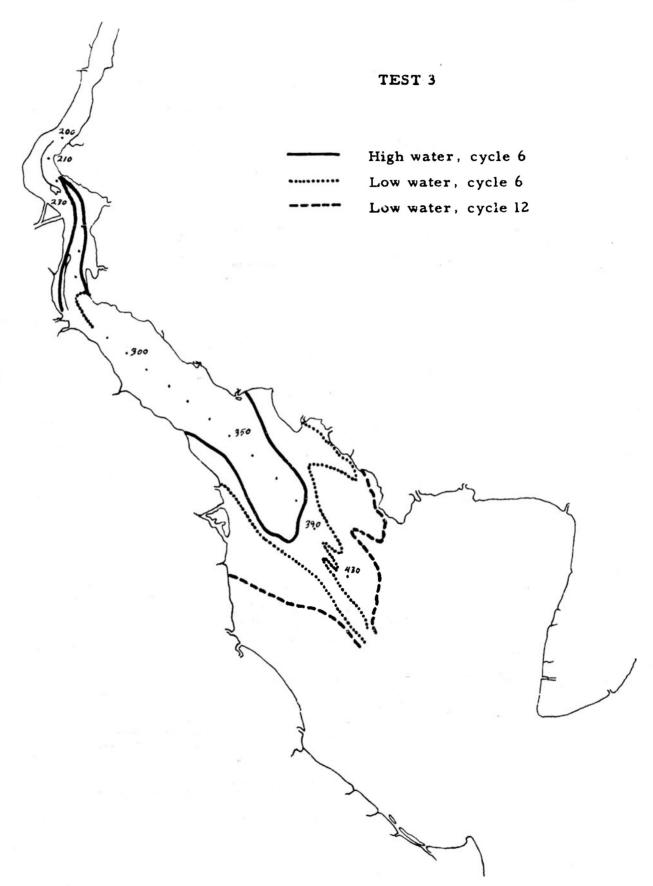
Visual Observations of the Spread of the Dye.
Tables of Observed Dye Concentrations.

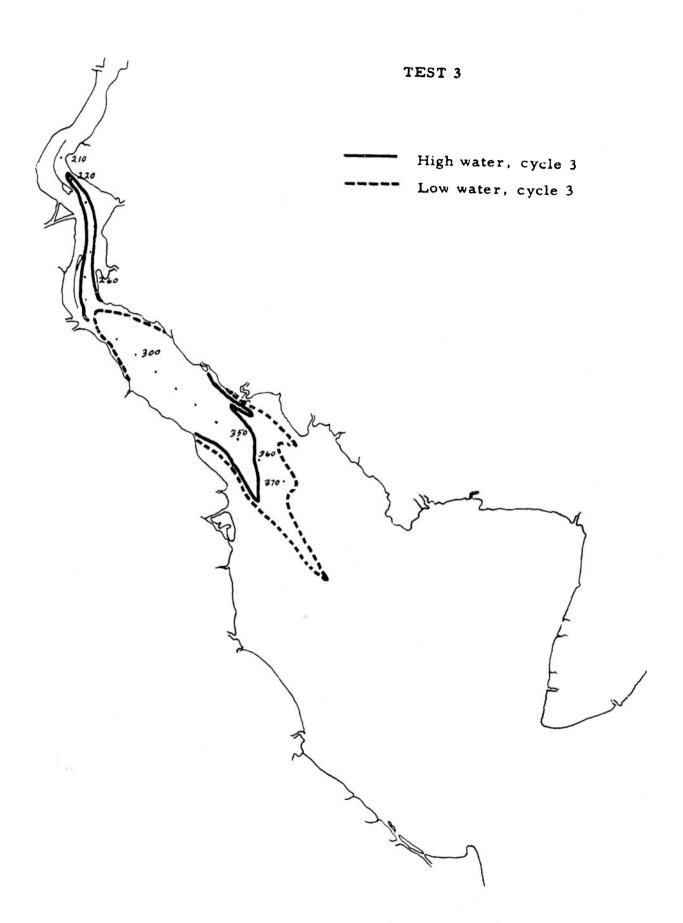
For Tests 3 through 7 the location of the visible limits of the dye was noted and charts were made during the tests. Notes were also made of any interesting features that were observed. These charts are presented in this appendix.

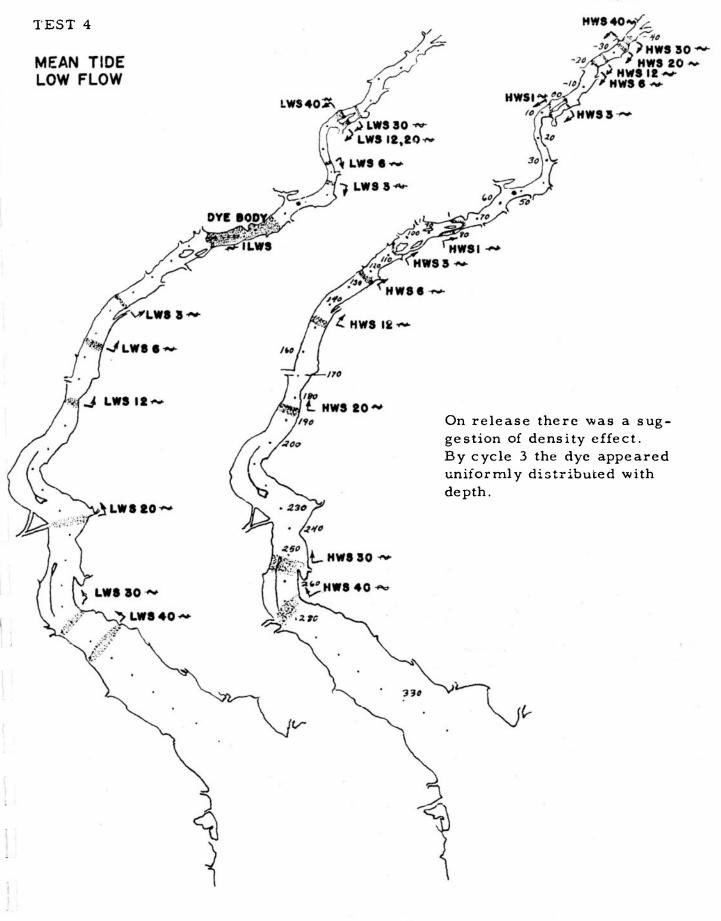
It was found that the limits of the dye concentration as determined visually corresponded to an average concentration of less than 0.1 ppm. This corresponds to more than a ten thousand fold dilution of the initial concentration.

This appendix also contains tables of the observed dye concentrations.

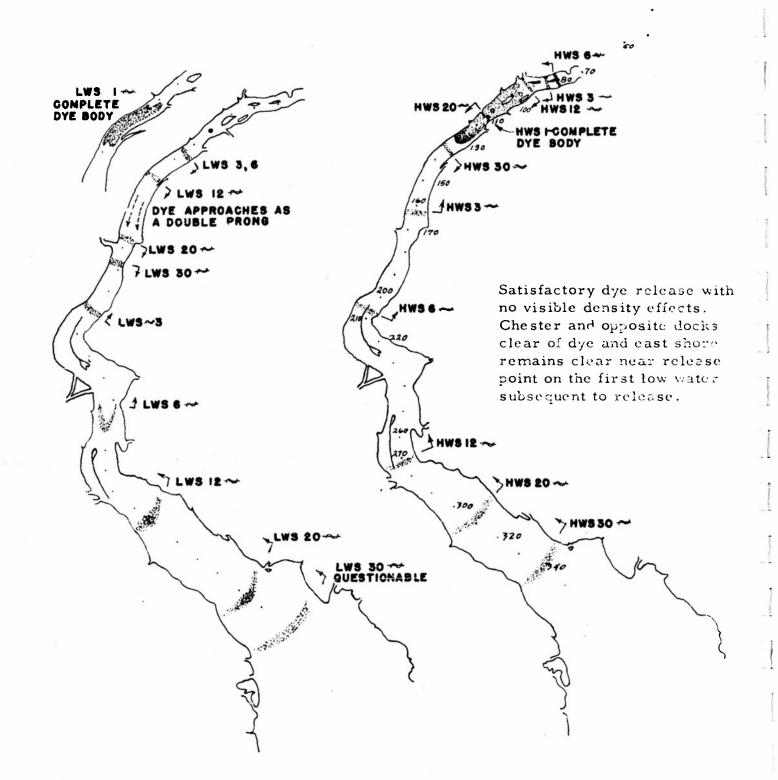


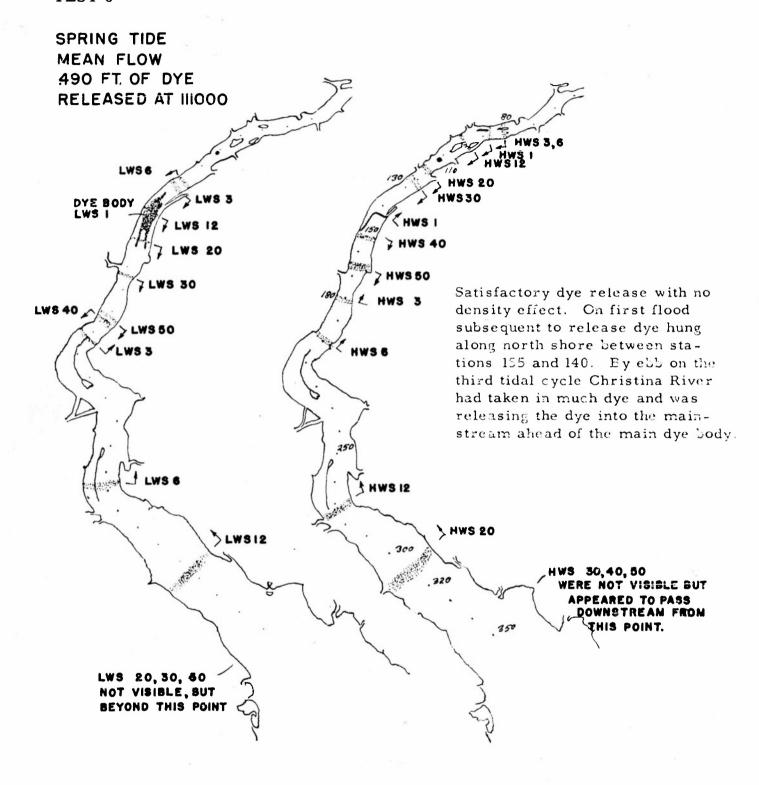




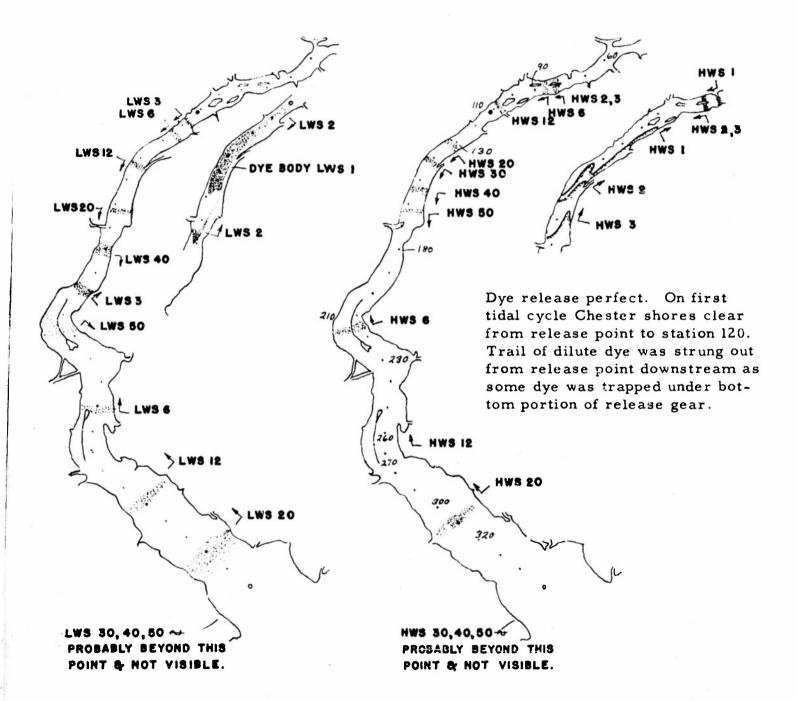


TEST 5
MEAN TIDE
MEAN FLOW
RELEASE AT CHESTER. III + 000





TEST 7



							
	Station	Depth	Conc. (ppm)		Station	Depth	(ppm)
	3 0	М	.084		-10	М	.006
TIDAL CYCLE 1	52.5	S M B	5.82 6.38 5.92		13	S M B	.023 .033 .016
					30	М	.071
7	13	S M B	.006 .037 .075		52.5	S M B	.469 .374 .456
TIDAL	30	м	.300	TIDAL	75	М	1.64
CYCLE 2	52.5	s	4.05	CYCLE 8	100	М	.171(?)
	M 4.10 B 4.21		127W*	S M B	1.53		
	75	М	1.22				1.11
	-10 52.5	M S	.006	*	127*	S 10 ft 20 ft 30 ft B	1.37 1.63 1.68 1.52 1.51
		M B	1.18 1.33		127E*	S B	1.19 1.33
TIDAL	127W*	S M B	.647 .587 .681		127EE*	М	1.22
CYCLE	127*	s 10 ft 20 ft 30 ft B	.612 .453 .702		160	М	.006
	127E*	S B	.447 .379				
	127EE*	м	.193				

^{*}Integrated sample over complete tidal cycle

TEST 1-A

	Station	Depth	Conc.		Station	Depth	Conc.
mrn ar	-10 13	M S M B	.000 .004 .000 .018		127*	S 10 20 30 B	.386 .426 .330 .381 .329
TIDAL CYCLE	30	М	.032		127E*	s	.381
16	52.5	S M	.143 .135	TIDAL CYCLE	·	В	•375
		В	.183	22 (cont'd)	127EE*	М	.396
	100	M	•559		160	М	•439
	127W*	S M	.687 .688		200 240	M	.194
		В	.753		240	S M	.083
	127*	S 10 20 30 B	.768 .853 .766 .867 .836		280	М	.072
	127 E *	S B	.678 .848		75	М	.011
	10000				100	М	.047
	127EE* 160	M M	.748 .362		127W*	s M	.098 .083
	200	M	.045			В	.111
	240	М	.054		127*	s 10	.107
	280	М	.016			20 30 B	.118 .122 .100
	52.5	S	.060	TIDAL CYCLE	127E*	S B	.100 .136
TIDAL		М	.075	28	127EE*	м	.128
CYCLE 22	75	М	.083		160	м	.418
	100	М	.185		200	S	.277
	127W*	S M B	.246 .253 .285			M B	.275 .277
		H	.20,		240	S M B	.278 .110 .073
					280	М	.042

TEST 1-A

	Station	Depth	Conc.		Station	Depth	Conc. (ppm)
	100	М	.031		127	М	.0 3 3
	127W*	S M B	.041		127EE	М	.031
	2	В	.051 .050		160	М	.113
	127*	S 10 20 30	.051 .051 .062 .046		200	S M B	.130 .156 .138
		В	.045		240	S M	.125 .127
TIDAL	1275*	S B	.032 .061			В	.097
CYCLE 34	127EE*	М	.045	TIDAL CYCLE 40	280	S M B	.113 .073 .097
	160	М	.300	(cont'd)	315	S	.038
	200	5 M B	.268 .227 .240			M B	.008 .026
	240		.185		350	M	.011
	_,,	S M B	.150	,	C&D Canal	М	.058
	280	М	.051	TIDAL	315	s	.045
	315	М	.011	CYCLE 49)1)	M B	.024
	127	M	.038		127	М	.004
	160	М	2ر1،	4	160	М	.031
TIDAL CYCLE	200	S M B	.202 .228 .227		200	S M B	.098 .093 .084
40	240	S M B	.157 .142 .125	TIDAL CYCLE	240	S M B	.074 .083 .077
	280	М	.056	52	280	S	:063
	315	М	.017			M B	.046 .058
					315	S M B	.037 .016 .021
					350	м	.011

TEST 1-A

	Station	<u>Depth</u>	Conc.		Station	<u>Depth</u>	Conc. (ppm)
	280	S M B	.107 .073 .078		200	S M B	.000 .024 .004
TIDAL CYCLE 5 ¹ 4 ¹ / ₂	315	S M B	.061 .032 .032		5#0	S M B	.012 .019 .024
	350	М	.032		280	S M	.016 .010
	160	М	.019	TIDAL CYCLE 64	280EE	B M	.004
	200	S M B	.011 .032 .017		280 e	М	.002
	240	S M B	.047 .075 .042		315	S M B	.000 .000
TIDAL CYCLE 58	280	S M B	.055 .044 .044		350	М	.000
	280EE	М	.081				
	315	S M B	.038 .027 .027	¥			
†	350	М	.019		a		

TEST 1

i							
	Station	<u>Depth</u>	Conc. (ppm)		<u>Statio</u> n	<u>Depth</u>	Conc. (ppm)
	54	S(c) M(c)	7.87 6.10		54	М	_ *
		B(c)	4.37		54R	M(c)	.618
	54R	M(c)	2.84		54L	M(c)	1.40
	54L	M(c)	6.02 *		60	M(c)	2.12
TIDAL	60	S(c) M(c)	6.30 8.45		70	М	- *
CYCLE		В	- *	TIDAL	80	M(c)	3.85
	70	S(c) M(c)	•795 •550	4	10	М	n.d.
		B(c)	1.60		0	M	n.d.
	80	S W	n.d.		90	M	- *
		м В	n.d.	595	100	M(c)	1.95*
	54	S(c)	3.05		110	M(c)	.120*
)+ 	M B(c)	- * .698		120	М	_ *
	54R	M(c)	1.63		54	М	- *
	54 L	M(c)	2.25*		54R	M(c)	.064
	60	S(c) M(c)	4.42 5.87		54L	M(c)	.205
TIDAL CYCLE	A	B(c)	4.94		60	M	_ *
2	70	S(c) M(c)	4.42 4.94		70	M(c)	.789
		B(c)	5.21	TIDAL	80	M(c)	1.43
	80	S(c) M(c)	.848 1.80	8	90	M(c)	2.42
	R	B(c)	2.14		100	М	_ *
	10	м	n.d.		110	M(c)	.888*
	90	s M	n.d. n.d.		120	M(c)	1.84
		M B(c)	n.d. .219		130	S(c)	•377 - *
	100	S(c)	.493			M B	- *
		M(c) D	.279 n.d.		140	s M	_ * _ *
*Evident	*Evident adsorption of dye on sample bottle n.d.						n.d.

Sample showed no visible evidence of dye--not run in photometer

				Г			
	Station	<u>Depth</u>	Conc. (ppm)		Station	Depth	Conc. (ppm)
TIDAL CYCLE 8 (cont'd)	150 160	M M	n.d.		130	S(c) M B(c)	•754 - * •620
	54	M(c)	.038		150	S M(c)	n.d. .493
	60 70	M M	- * - *		160	B(c) S(c)	.013
	80	М	_ *			M B(c)	- * •447
	90	М	_ *	TIDAL	170	М	- *
TIDAL	100 110	М(с) М(с)	.675 1.38	16 (cont'd)	180	S M B	n.d. - * n.d.
CYCLE 12	120	M(c)	1.90		190	M(c)	.115
	130	S	_ *		200	M(c)	.085
	*	M(c) B(c)	1.23 1.60		210	м	n.d.
	140	S(c) M	•990 - *		220	м	n.d.
		B(c)	•9 7 5		240	М	n.d.
	150	S(c) M(c) B(c)	1.10 .813 .497		54	М	.020
	160	М	- *		60 70	M M	n.d.
	54	М	n.d.		90	М	n.d.
(7	60	М	n.d.		100	м	r.d.
	70	М	n.d.	TIDAL	110	M	n.d.
TIDAL CYCLE	90	M(c)	.175	22	120	M(c)	.060
16	100	M	<u>*</u> *		130	M(c) B	.020 n.d.
	110	M M(c)	- * .668		140	S M(c) B	n.d. .000 n.d.

	Sta+*.on	Depth	Conc. (ppm)		Station	Depth	Conc. (ppm)
	-150	S M(c) B	n.d. .169 n.d.		170 180	M(c) S(c)	.279 .175
	160	S B	- * - *		190	M(c) B(c) M	.175 .432 .386
	170	M(c)	.352		200	S	n.d.
TIDAL CYCLE	180	S M(c) B(c)	- * .620 .352			M(c) B	.041 n.d.
22 (cont'd)	190	M(c)	.475	27	220	S M(c) B	n.d. .210 n.d.
	200	S M(c) B	n.d. .000 n.d.	TIDAL CYCLE 28 (cont'd)	240	S(c) M(c) B(c)	.017 .056 .056
	220	S M(c) B	n.d. .100 n.d.	(cons u)	260	S M B	n.d. n.d. n.d.
	240	M	n.d.		280	S	n.d.
	260	М	n.d.		200	M B	n.d. n.d.
	280	М	n.d.		300		n.d.
	90	М	n.d.		900	S M B	n.d. n.d.
	100	М	n.d.				
	110	M(c)	.013				
TIDAL CYCLE 28	130	S M B	n.d. n.d. n.d.				2
	140	M(c)	.026				
	150	S M(c) B	n.d. .026 n.d.				
	160	S(c) B(c)	.041 .065				

	7		
	Station	<u>Depth</u>	Conc. (ppm)
	90	М	n.d.
	100	М	n.d.
	110	М	n.d.
	130	S M B	n.d. n.d. n.d.
	140	м	n.d.
	150	S M(c) B	n.d. .020 n.d.
	160	S M(c) B(c)	n d. .055 .013
TIDAL	170	M(c)	.026
CYCLE 34	180	S M B(c)	n.d. - * .017
	190	М	_ *
2	200	S M B	n.d. n.d. n.d.
	220	S M(c) B	n.d. .130 n.d.
	240	S M B	n.d. n.d. n.d.
	260	S M B	n.d. n.d. n.d.
	280	S M B	n.d. n.d. n.d.

	Station	Depth	Conc. (ppm)
TIDAL	300	S M(c) B	n.d. .020 n.d.
CYCLE 34	3 20	М	n.đ.

	Station	Depth	Conc. (ppm)	14,	Station	<u>Depth</u>	Conc. (ppm)
	100	M(b)	.008	4	100	M(b)	.∞₀
	110	м(ъ)	.000	. (18)	110	M(b)	.001
	130	S(b) M(b) B(b)	.000 .055 .001		130	S M(b) B(b)	n.d. .000 .000
	150	M(b) B(b)	.000		150	М	n.d.
	160	S M B(b)	n.d. n.d.	*1	160	S M(b) B	n.d. .009 n.d.
	180	S(b) M(b) B(b)	.0 00 .008 .004	TIDAL CYCLE 46	180	S M(b) B	n.d. .003 n.d.
	200	S	n.d.		200	M(c)	.026
	200	М В(с)	n.d. .070		220	S M B	n.d. n.d. n.ā.
TIDAL	220	S(c) M(c) B(c)	.110 .060 .060		240	M(b) B(b)	.014 .015
CYCLE 40	240	S	n.d. .041		260	М(ъ)	.045
į		M(c) B(c)	.000		280	M(b)	.001
	260	S M(c) B	n.d. .001 n.d.		300	S(b) M(b) B	.020 .000 .000
	280	S	n.d. n.d.		320	s	n.d.
		M B	n.d.		350	М(b) В	.001 n.d.
	300	S(c) M B(c)	.041 n.d. .013			Б	n.u.
	320	S M B	n.d. n.d. n.d.				-
	350	М	n.d.				

TEST 1

	Station	Depth	Conc.
	100	M(b)	.000
	110	M(b)	.000
	130	S(b) M(b) B(b)	.000 .000
	150	м(ъ)	.000
an an	160	S(b) M(b) B(b)	.000
	180	s M(b) B	n.d. .000 n.d.
TIDAL	200	S(b) B	.003
52 CYCLE	220	S(b) M(b) B(b)	.009 .017 .042
	240	S(b) M(b)	.054 .019
y	260	S(b) M(b) B(b)	.010 .008 .000
	280	S(b) M(b) B(b)	.022 .026 .016
A 4	3 00	S(b) M(b) B(b)	.010 .007 .014
	32 0 -	S(b) M(b)	.005 .006
	350	S M B	.007 .000 .000

	Station	<u>Depth</u>	Conc. (ppm)
TIDAL	300	М	.045
CYCLE 56	320	М	.031
	200	М	.005
	550	М	.022
TIDAL	240	М	.044
CYCLE 58	260	М	.049
	280	м	.035
	350	М	.023

		· · · · · · · · · · · · · · · · · · ·	
	Station	<u>Depth</u>	Conc.
	220	S M B	n.d. n.d. n.d.
	240	S M B	n.d. n.d. .022
	260	S M B	.215 .605 .540
	280	S M B	.740 .870 .990
TIDAL	300	S M B	1.24 1.25 1.23
6	300E	М	.885
	300W	М	2.73
	320	S M B	.955 1.09 1.01
	320E	М	.855
	320W	М	2.04
	340	S M B	1.40 .800 .456
	350	S M B	.780 .334 - *
	350E	М	.100
	350W	м	.930
	370	S M B	.165 .054 .039

	Station	Depth	Conc.
TIDAL CYCLE 6	390	S M B	n.d. n.d. n.d.
	390E	М	n.d.
	390W	М	n.d.
(cont'd)	410	S M B	n.d. n.d. .005

TEST 2

	Stati on	Depth	Conc.
	180	M	n.d.
	200	М	n.d.
	220	S M B	n.d. .039 .063
	240	S M B	.02 ⁾ ! .103 .063
	260	S M B	.073 .170 .140
	280	S M B	.186 .230 .158
TIDAL CYCLE 24	300	S M B	.175 .347 .321
	300 e	М	.460
p.	300W	М	.469
	320	S M B	.302 .386 .315
7.	320E	М	.442
	320W	М	.406
€	340	М	-347
	350	S M B	.180 .066 .117
	350 e	M	.370

	Station	<u>Depth</u>	Conc. (ppm)
	350W	M	.460
	370	S M B	.194 .153 .238
	390	M B	.199 .162 .190
TIDAL	390E	М	.080
CYCLE 24	390W	М	.158
(cont'd	410	S M B	.085 .054 .054
2	430	S M B	.028 n.d. n.d.
	430E	М	n.ā.
	430W	М	n.d.
	_		

	Station	Depth	Conc. (ppm)
	180	М	n.d.
	200	М	.020
	220	S M B	.024 .038 .024
	240	S M B	.015 .058 .000
	260	S M B	.000 .063 .149
	280	s M	.126 .152
TIDAL CYCLE	300	s M B	.134 .135 .135
36	30 0E	М	.185
	300W	М	.213
	320	s M B	.315 .285 .208
	320E	М	.239
	320W	М	.410
	340	М	.267
	350	S M B	.000 .267 .135
	350 E	М	,217
	350W	M	.140
	370	S M B	.081 .117 .135

	Station	Depth	Conc.
TIDAL CYCLE 36	390	S M B	.117 .073 .063
	39Œ	М	.20 S
	390W	М	.103
	410	S M B	n.d. .032 n.d.
	430	S M B	n.d. n.d. .066
	430 E	М	.050
	430W	М	n.d.

TEST 2

		T	
	Station	Depth	Conc.
	180	М	n.d.
	200	M	n.đ.
	220	S M B	n.d. n.d. .020
	240	S M B	.020 .020 n.d.
	260	S M B	.040 .027 .040
	300	S M B	.038 .094 .038
TIDAL CYCLE	300E	М	.226
42	300W	М	.180
	320	S M B	.1.45 .099 .095
	320E	M	.235
	320W	М	.239
	340	М	.185
	350	S M B	.066 .099 .084
# 1	350 E	M	.238
	350W	M	130
	370	S M B	.080 .046 .099

	Station	<u>Depth</u>	Conc.
	390	S M B	.063 .054 .049
	390E	М	.172
	390W	М	.080
TIDAL CYCIE 42 (cont'd)	410	S M B	.046 .032 .040
	430	S M B	n.d. .020 n.d.
	430E	М	.054
	430W	М	n.d.

TEST 3

	<u>Station</u>	<u>Depth</u>	Conc. (ppm)	5	Station	Depth	Conc. (ppm)
	260	S M B	.068 .265 .905		260	S M B	.347 1.37 1.44
22.5	280	S M B	.35° 2.55 4.02		280	S M B	.989 2.18 1.67
	280 e	М	.615		280E	М	1.24
	280W	М	1.12		280W	м	1.99
	300	S M B	2.07 8.00 3.84		300	S M B	2.08 3.25 2.74
TIDAL	300E	м	.00ರ	TIDAL	300E		1.85
CYCLE	300W	М	.450	3	300W		2.34
	320	S M B	.322 .265 .111		320	S M B	2.75 1.46 1.09
	320E	М	.033		320E	М	1.95
F #	320W	М	•377		320W	М	3 . 35
	330	S M B	.127 .029 .037		330	S M B	1.28 .905 .320
	350	S M B	.016 .033 .143		350	S M B	.278 .074 .017
70	350E	M	•044		350E	М	.033
	350W	M	.037		350W	М	.231
						4	

	Station	Depth	Conc. (ppm)		Station	Depth	Conc. (ppm)
	220	М	.049		220	. М	.017
	240	S M B	.100 .171 .201		240	S M B	.074 .143 .099
	260	М	.650		280	S M	.465 .420
	280	S M	.762 1.28		300	B M	.295
		В	1.35				.889
	280E	M M	1.13		320	S M B	.869
	300	M	1.38		320E	М	.873
	30 0 E	M	1.34		320W	м	.858
	300W	М	1.42		330	М	•975
TIDAL CYCLE	320	S M B	1.38 1.11 .868	TIDAL	350	S M B	.830 .619 .3 8 2
6	320 E	М	1.55	12	350E	М	.566
	320W	М	1.63		350W	М	.842
	330	S	1.44		390	М	.205
		M B	1.08 .713		390E	М	.119
	350	S	1.00		390W	М	.219
		M B	.49 <u>1</u> .320		430	S M	.099
	350 E	М	.241			В	.029
	350W	М	1.16		430E	М	.021
	370	М	.075		430W	М	.017
	390	S M B	.045 .016 .020		25		
	390E	М	.043				
	390W	м	.143				
			1	I			

.041

M

410

	Station	<u>Depth</u>	Conc. (ppm)		Station	Depth	Conc. (ppm)
	220	М	.017		220	М	•00jt
	240	S M B	.074 .041 .043		240	S M B	.043 .017 .016
	280	S M B	.185 .250 .279		280	S M B	.103 .214 .214
	300	М	.484		300	М	. 250
	320	S M B	.365 .400 .410		320	s M B	.285 .289 . 3 00
	320E	М	.471		320E	м	.275
	320W	М	.409	TIDAL	320W	м	.268
TIDAL	330	М	.510	CYCLE 30	330	М	.250
20	350	S M B	.360 .489 .435		350	S M B	.300 .265 .265
	350E	М	.521		350E	М	.345
	350W	M	.571		350W	М	.325
	390	M	.233		390	М	.176
	390E	М	.280		390E	M I	.302
	390W	М	.301		390W	М	•339
	430	S M B	.099 .068 .070		430	S M B	.119 .070 .053
	430E	М	.046		430E	М	.127
	430W	М	.020		430W	М	.049

	Station	Depth	Conc.
	220	М	.005
	240	S M B	.004 .004 .008
	280	S M	.061 .083
	300	М	.184
	320	S M B	.159 .176 .131
	320E	М	.196
	320W	M	.168
	330	М	.176
TIDAL CYCLE 40	350	S M B	.168 .180 .151
	350E	М	.188
	350W	M	.155
	390	М	.111
II	390E	М	.214
	390W	М	.129
	430	S M B	.049 .082 .021
6	430E	М	.100
	430W	М	.070

	Station	Depth	Conc. (ppm)
	280	S M B	.053 .059 .083
10	300	М	.090
	320	S M B	.090 .100 .090
	320E	М	.119
	320W	М	.115
	330	М	.089
TIDAL	350	S M B	.107 .100 .083
CYCLE 50	350E	М	.127
	350W	М	.131
	390	М	.041
	390E	М	.143
	390W	М	.107
	430	S M B	.033 .016 .012
	430E	М	.090
	430W	М	.04.1
·	1	<u> </u>	

į	Station	<u>Depth</u>	Conc. (ppm)
	20	S M B	.138 .094 .106
;	30	М	.271
	40	S M B	2.21 1.56 .7 7 0
	54	S M B	>10 >10 >10
TIDAL	54R	М	8.60
CYCLE	54 L	М	8.65
	60	S M B	2.98 2.98 3.50
	70	М	.138
	80	S M B	.028 .086 .061
	1		

9	Station	Depth	Cone. (ppm)
	10	M	.280
	20	S M B	.451 .505 .538
!	30	М	1.50
	40	S M B	1.95 2.26 1.84
TIDAL CYCLE 3	54	S M B	5.28 4.64 4.55
)	54R	М	3.50
	54L	М	5.91
	60	S M B	5.12 3.71 4.08
	70	м	2.04
	80	S M B	.840 .912 1.07
	100	S M B	.170 .199 .178

TEST 4

	Station	<u>Depth</u>	Conc.		Station	<u>Depth</u>	Conc.
	20	S M B	.670 .638 .530		-30 -10	M M	.069 .178
	40	S M B	1.79 1.74 1.63	54.0 ST	10 20	M	.340 .451
	54R	M	2.09		20	S M B	.490 .495
	54L	М	2.98		40	M	1.11
TIDAL CYCLE 6	60	M B	2.56 2.15	TIDAL CYCLE	54	S M B	1.57 1.42 1.35
	70	M	2.48	12	80	м	1.90
	80	s M B	1.61 1.82 1.78		100	S M B	1.32 1.24 1.32
	90	М	1.29		120	S	.510
	100	S M B	1.02 1.10 .711		140	B S B	.749 .061 .089
	120	S M B	.203 .170 .411			д	.009

TEST 4

	Station	Depth	Conc.		Station	<u>Depth</u>	Conc.
	-30	М	.073		-50	М	.056
	-10	М	.21.4		-30	М	.073
	10	M	.311	İ	-10	М	.130
	20	S M	.251 .271		10	- M	.230
		В	.195		20	М	.166
	40	S M	.540 .468		30	М	.300
		В	.389	TIDAL	40	s M	.283 .259
TIDAL CYCLE	60	s M	.800 .730	30		В	.292
20		В	.745		60	.s M	•591 •485
	80	s M	1.35 1.36			3	.440
i		В	1.20	ļ	80	S M	.840 .800
	100	s M	1.36 1.20			В	.840
·		В	1.29		100	s M	.94 3 .928
	320	s M	.790 .705			В	1.04
-		В	.699		120	M	•741
	140	s M	.331 .352		140	s M	.588 50 7
		M B	•337		**	В	• 534
	160	S M	.315 .126		160	M	.263
		В	.126		180	s M	.214 .190
=	180	s M	.085 .053			В	.170
	20	В	.081		200	М	.118
	200	М	.065		220	М	.101
					240	M	.073
					260	M	.065
		55					
				ļ			

		···
Station	Depth	Conc. (ppm)
-50	М	.036
- 30	М	.053
-10	М	.069
10	М	.118
20	М	.065
60	S M B	.304 .288 .292
80	S M B	.255 .360 .449
100	S M B	.540 .538 .469
120	М	.630
140	S M B	.544 .545 .538
160	М	.364
180	S M B	.255 .235 .248
200	S M B	.000 .259 .122
220	м	.153
240	М	.089
260	М	.060
	-50 -30 -10 10 20 60 80 100 120 140 160 180 200 220 240	-50 M -30 M -10 M 20 M 20 M 60 S M B 100 S M B 120 M 140 S M B 160 M 180 S M B 200 S M B 200 M

	Station	Depth	Conc. (ppm)
TIDAL	280	М	.056
CYCIE 40 (cont'd)	300	M	.056

	Station	Depth	Conc.
	80	М	.034
	90	S M B	.184 .093 .034
	100	s M	.223 .083
TIDAL	110	S M B	3.35 3.15 2.18
1	120	S M B	7.12 7.02 3.82
	130	S M B	.246 .367 1.03
	140	М	.026

	Station	Depth	Conc. (ppm)
	70	М	.118
	80	М	. 398
54 Y	90	S M B	.078 .165 .093
	100	s M B	.219 .181 .205
	110	S M B	1.35 1.23 1.22
TIDAL CYCLE 3	120	S M B	3.82 4.15 3.65
	130	S M B	3.86 4.16 4.19
	140	S M B	1.63 1.66 1.58
	150	S M B	.336 .398 .332
	160	М	.098

TEST 5

·						
	Station	Depth	Conc. (ppm)			<u>s</u>
	70	М	.000			8
	80	М	.012			9
	90	S	.066			1
		M B	.072 .062	-0	1	. 1
	110	S M B	• 39 4 • 367 •348			1
	130	S M B	.000 2.30 2.30		TIDAL CYCLE	1
TIDAL CYCIE 6	140	S M B	2.38 2.58 2.51		12	נ
	150	S M B	1.97 1.86 1.70]
	160	S	1.08 .885			2
		M B	.900			2
	170	M	.778			2
	190	М	.102			
					-	- 1
L	1		i	1	L	L

	Station	<u>Depth</u>	Conc. (ppm)
	80	М	.026
	90	М	.012
	100	М	.053
	130	S M B	.493 .508 .466
	140	М	•797
TIDAL	150	S M B	1.22 1.32 1.30
CYCLE 12	160	М	1.45
	170	S M B	1.17 1.21 1.30
	190	М	.645
	210	М	.263
	230	М	.259
	250	м	.078
	j.		4

	Station	Depth	Conc.
	90	М	.008
	110	n	.026
	130	8 M B	.152 .120 .120
TIDAL CYCLE	150	S M B	.513 .304 .348
20	170	s M B	.693 .680 .645
	190	M	.600
	210	М	.546
	230	М	.453
	250	М	.278
o.	270	М	.098

	Station	Depth	Conc.	
	110	М	.026	
	130	S M B	.048 .043 .066	
	150	М	.098	
TIDAL CYCLE 30	170	S M B	.223 .228 .233	
	190	S M B	.256 .308 .304	
	210	S M B	.300 .358 .381	
	230	м	•394	
	250	S M B	.332 .286 .283	
	270	S M B	.283 .219 .173	
	290	м	.242	
	31 0	М	.102	

			
	Station	D pth	Conc.
	80	М	.023
***	90	S M B	.000 .071 .071
	100	S M B	.079 .083 .179
TIDAL CYCLE	110	S M B	2.96 2.27 3.40
1	120	S M B	5.60 5.97 6.45
	130	S M B	•583 •535 •463
	140	М	.052
į			

	Station	Depth	Conc.
	70	М	.012
	80	М	.150
	90	S M B	.047 .054 .058
TIDAL CYCLE 3	100	S M B	.063 .131 .219
	110	S M B	1.10 .950 1.09
	120	S M B	3.24 3.20 3.24
	130	S M B	3.22 3.26 3.10
	140	s M B	1.85 1.64 1.61
	150	S M B	•553 •435 •307
	160	М	.148
	170	M	.135

	Station	Depth	Conc. (ppm)		Station	<u>Depth</u>	Conc. (ppm)
	70	М	.052		80	М	.003
E 2	80	М	.071		90	М	.012
	90	S	.043 .098		110	М	.071
		M B	.063		130	S M	•395 •419
	110	S M	.035 .207			В	.415
		В	.372		140	М	.657
	130	S M B	1.97 1.97 1.90		150	S M B	.926 .483 .830
TIDAL	140	S	2.20	TIDAL	160	м	.981
CYCLE 6		M B	1.98 1.94	12	170	S M	.774 .457(?)
	150	S M B	1.88 1.95 1.75		190	B S M	.750 .527 .423 .488
	160	S M	•799 •857		4	В	
	ě.	В	1.05		210	S M	.367 .251 .248
	170	S M	1.15(?) .387			В	
		В	.352		230	M	.,074
3	190	М	.111		250	М	.039
312	210	M	.027		270	М	.030

	GL-Adam	Depth	Conc.
	Station	Depth	(ppm)
	90	М	.007
	110	м	.030
	130	S M B	.079 .047 .103
	150	s M B	.183 .203 .215
	170	S M B	.363 .381 .387
TIDAL CYCLE	190	s M B	.483 .500 .478
20	210	S M B	.443 .463 .427
	230	М	.358
	250	М	.263
	8	£4	
	270	S M B	.299 .235 .292
	290	М	.167
	310	М	.079

	Station	Depth	Conc.
	110	М	.000
	130	S M B	.030 .019 .054
	150	М	.091
TIDAL CYCLE 30	170	S M B	.087 .154 .188
	190	М	.243
	210	S M B	.299 .295 .318
	230	М	.332
	250	М	.303
	290	М	.183
	310	М	.079
	330	М	.054
	350	М	.047

Programme de

Constitution 4

TEST 6

	Station	Depth	Conc.		Station	Depth	Conc. (ppm)
	130	М	.030		150	М	.000
	150	M	.030		170	S M	.047 .035
	170	S M B	.035 .083 .103		190	B M	.054
	190	М	.119		210	S M	.103 .115 .087
	210	S M B	.171 .191 .163		230	В М	.119
TIDAL CYCLE	230	М	.199	TIDAL CYCLE	250	s M	.143 .167
40	250	S M B	.235 .183 .243	50	270	В м	.143 .187
	270	М	.188		290	S M	.139 .111
	290	S M	.188		7 70	В м	.111
		В	.143		310		.071
. 36	310	М	.087		330	М	et a
K1	330	М —	.083		350	М	.063
	350	М	.054		390	М	۰054
	430	М	.019		430	М	.027

	Station	Depth	Conc.
	80	М	.013
	90	S M B	.050 .079 .058
	100	S M B	.097 .109 .025
TIDAL	110	S M B	4.45 4.28 3.6 1
1	120	8 M B	6.25 5.62 10.0
	130	S M B	.280 .298 .433
	140	М	.088

·	Station	Depth	Conc. (ppm)
	70	М	.013
	80	М	.050
	90	S M B	.125 .071 .050
	100	S M B	.092 .147 .213
l	110	S M B	1.15 1.10 1.59
TIDAL GYCLE 3	120	S M B	4.06 3.65 3.64
	130	S M B	4.38 3.85 3.87
	140	S M B	1.05 1.10 1.03
	150	S M B	.363 .259 .188
	160	М	.117
			1

	Station	Depth	Conc.		Station	Depth	Conc. (ppm)
	70	М	.003		80	М	.003
	80	M	.029	8	90	М	.003
	90	S M	.100		110	М	.021
		· · · · · · · · · · · · · · · · · · ·	.104		130	s M	.417 .500
	110	s M	.255 .008			В	.525
 		В	.275		140	М	.625
	130	S M	1.87 1.99	TIDAL	150	S M	.8 03 .950
TIDAL		В	1.91	12		В	1.05
CYCLE 6	140	S M	2.20 2.36		160	М	1.18
		В	2.22		170	S M	.741 .988
	150	S M	1.96 1.72	l		В	1.00
		В	1.60		190	М	•592
	160	s M	1.22 .810		210	М	.272
	4	В	.961		230	М	.138
	170	м	-535		250	М	.400
	190	м	.150				
	210	М	.050				

		Station	Depth	Conc.	
		90	M	.037	
		110	M	.008	
	*	130	S M B	.075 .067 .088	
	20	150	s M B	.204 .259 .273	
TIDAL CYCLE		170	S M B	.363 .433 .550	
20	•	190	S M B	.467 .483 .471	-
		210	М	.467	
		230	М	.284	
	10	250	М	.204	
XX		270	М	.142	
		290	М	.130	
		310	М	.075	

	Station	<u>Depth</u>	Conc. (ppm)
	130	S M B	.047 .008 .050
	150	М	,05 8
	170	S M B	.138 .134 .100
	190	М	.213
TIDAL CYCLE	210	S M B	.242 .242 .259
30	230	м	.273
	250	М	.250
	270	М	.192
	290	М	.150
	310	М	.104
	330	М	.050
	350	М	.033
	Onthe	, =	
		197	ļ

TEST 7

	Station	Depth	Conc. (ppm)			Station	Depth	Conc.
	130	М	.029			150	М	.025
	150	М	.033	İ		170	s M	.033 .033
	170	S M B	.050 .047 .067			190	B M	.025
	190	M	.121			210	S M	.063 .054
	210	S M B	.154 .176 .172			230	B	.083
	230	М	.142			250	S M	.121
TIDAL CYCLE 40	250	S M B	.234 .204 .234		TIDAL CYCLE 50	270	B M	.142 .147
	270	М	.157			290	М	.109
	290	S M B	.225 .142 .200			310	S M B	.092 .067 .079
	310	м	.104			330	М	.071
	330	м	.079			350	M	.063
	350	М	.067			390	M	.047
	430	М	.042		`	430	М	.042
				Į				

APPENDIX III

Correction of Observed Concentrations.

Tables of Corrected Concentration.

The high adsorption of the methylene blue on any surface with which it came in contact -- the plaster and concrete of the model, dust, sample jars, and even the photometer cells -- caused considerable difficulty. It was found in laboratory tests that the adsorption on the model itself could be materially reduced by painting it with a clear plastic paint and this was done between Test l-A and the rest of the tests. The residual loss on the model, together with the losses on other surfaces, was compensated by a correction factor calculated in the following way.

If no dye were lost then the integral of the dye concentration, expressed as weight per unit volume, over the entire volume of the model would equal the weight of the dye introduced. The difference between this integral and the original weight of dye is therefore a measure of the dye lost by adsorption. The ratio of the integral to the weight of dye introduced may serve as a correction factor for changing observed concentration to corrected concentration which is an estimate of the value the observed concentration would have shown had adsorption not been present.

Consideration of the stations where samples were taken across the estuary showed that after the first one or two tidal cycles the mean sectional concentration could be estimated from the mid-channel samples. Using the mid-channel observations, concentration was plotted against longitudinal distance and a smooth curve drawn through the points. An

example of such a curve is shown in Figure A-III-1 where the open circles denote concentrations estimated from single samples and the solid circles denote concentrations estimated from an average of samples at surface, middepth, and bottom. In drawing the smoothed curves more weight was given to these latter points when a departure from the actual estimates seemed necessary.

From the smoothed curves the concentration at each 10,000 foot station was read. Each concentration was then multiplied by the appropriate mean cross sectional area and the integration was carried out numerically. The ratio of the total dye represented by the distribution curve to the amount of dye originally introduced was formed. This ratio was computed for each tidal cycle and plotted as a function of time. A smooth curve through these points provided the correction factor.

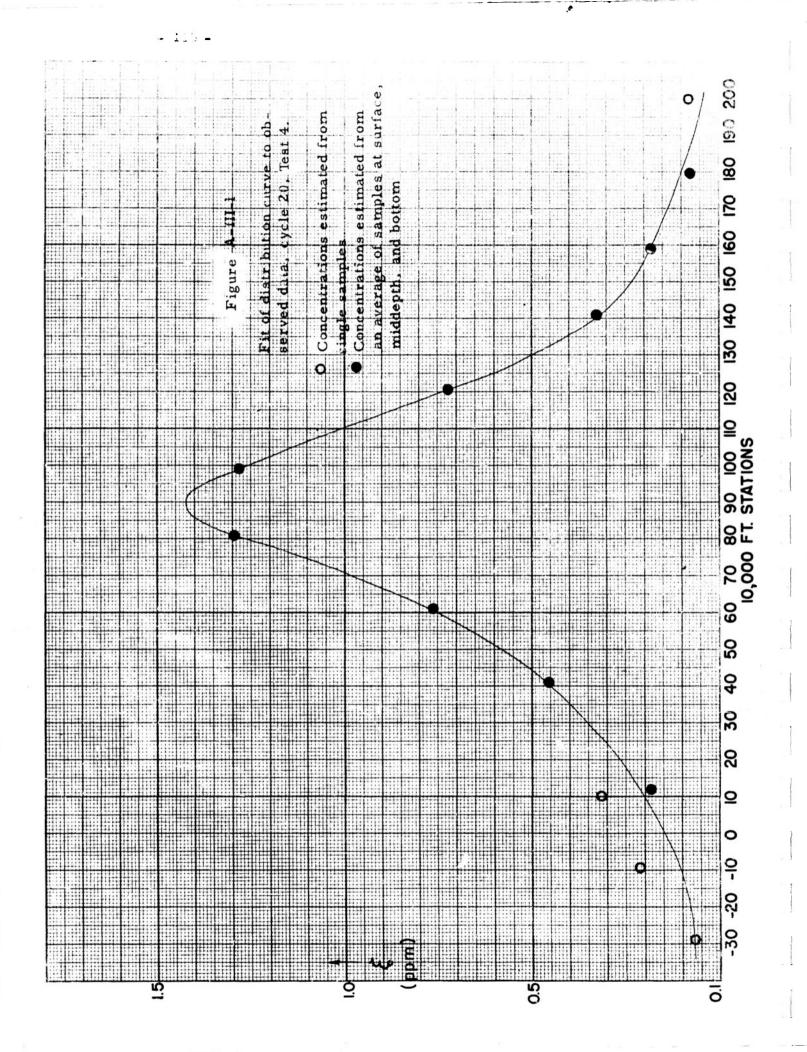
Four correction curves of this sort were used, one for Test 1-A, one for Tests 1 and 4, one for Tests 2 and 3, and one for Tests 5, 6, and 7.

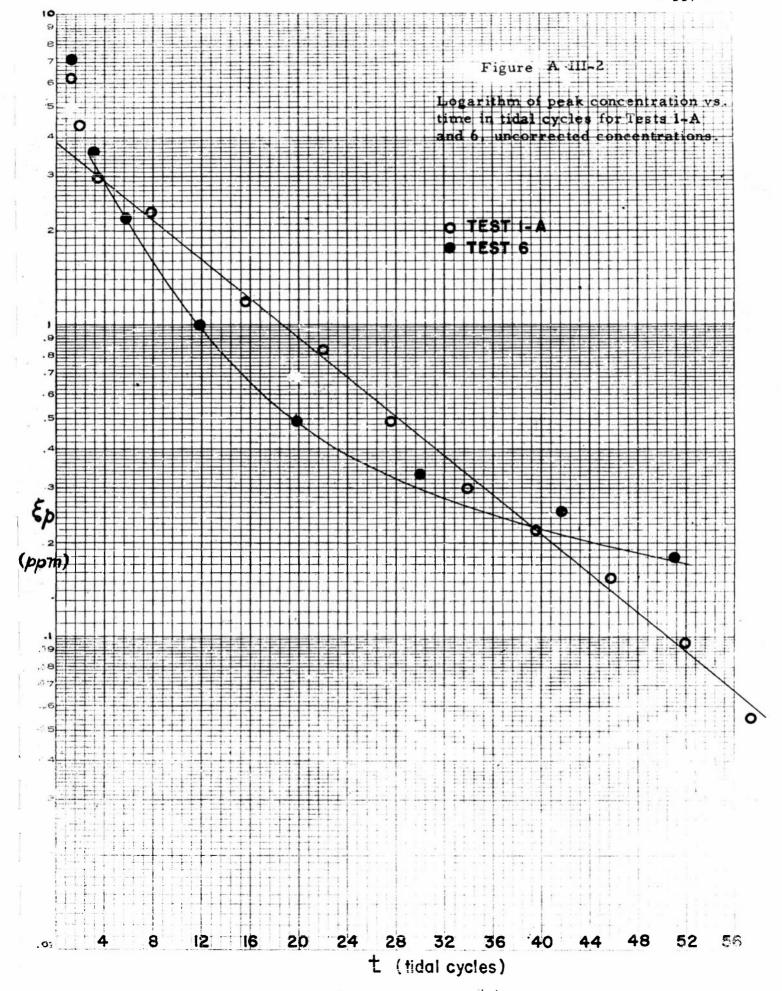
As was to be expected, the correction factors for Test 1-A, run before the model had been painted, were significantly higher than those for Test 1, run after the model had been painted, although the tests were otherwise the same.

The data analyzed in the text of this report have all been corrected in this way. When the difference in the nature of the decay during the later tidal periods between the first five and the last three tests was noted the question arose as to whether the difference was real or an artifact of the

data are used, although with somewhat difference still remains if raw data are used, although with somewhat different values for the logarithmic slopes, it seems unlikely that the difference is a result of the applied correction. Figure A-III-2 shows the uncorrected data for Tests l-A and 6 as an example. For Test l-A the logarithm of concentration versus time is still approximately linear while for Test 6 the straight line is still not suitable. Figure A-III-2 should be compared with Figure 24 which shows the same plot for the corrected data from the same two tests.

The table in this appendix gives the corrected concentrations for each tidal cycle by 10,000 foot stations.





TEST 1-A - DELAWARE RIVER

			×			
	Station	Corrected			Station	Corrected
TIDAL CYCLE 1	20 30 40 50 53.5* 60 70 80	0.03 0.17 1.90 10.30 10.56 6.09 0.52 0.03		TIDAL CYCIE 8 (cont'd)	100 110 120 130 140 150 160	3.02 1.89 0.80 0.23 0.08 0.05 0.03
TIDAL CYCLE 2	10 20 30 40 50 57.5* 60 70 80 90	0.05 0.20 0.47 1.49 5.57 6.83 6.77 3.31 1.21 0.31		TIDAL	20 30 40 50 60 70 80 90 100 110 120 128*	0.02 0.05 0.13 0.24 0.34 0.45 0.45 0.74 0.96 1.43 1.81
TIDAL CYCLE 4	10 20 30 40 50 60 70 75* 80 90 100	0.02 0.14 0.38 0.96 1.79 2.91 4.24 4.48 4.28 2.66 1.07 0.24		CYCLE 16	130 140 150 160 170 180 190 200 210 220 240 250 260 270 280	1.93 1.60 1.08 0.59 0.34 0.24 0.13 0.10 0.08 0.08 0.08 0.07 0.05 0.05 0.03
TIDAL CYCLE 8	20 30 40 50 60 70 80 90 92.5*	0.03 0.11 0.27 0.57 1.19 2.05 2.97 3.34 3.35		TIDAL CYCLE 22	40 50 60 70 80 90 100	0.05 0.11 0.13 0.14 0.18 0.23 0.32

* Location peak concentration

TEST 1-A

	Station	Corrected Conc.
TIDAL CYCLE 22 (cont'd)	110 120 130 140* 150 160 170 180 190 200 210 220 230 240 250 260 270	0.50 0.85 1.31 1.48 1.30 0.77 0.59 0.50 0.43 0.36 0.27 0.27 0.27 0.20 0.18 0.14 0.11 0.09
TIDAL CYCLE 28	80 90 100 110 130 140 153* 160 170 180 200 210 220 230 240 250 260 270 280	0.04 0.10 0.11 0.25 0.38 0.57 0.95 0.95 0.65 0.57 0.48 0.34 0.30 0.25 0.17 0.11

	Station	Corrected Conc.
TIDAL CYCLE 34	100 110 120 130 140 150 160 170* 130 290 210 220 230 240 250 260 270 280 290 300 310	0.08 0.15 0.21 0.34 0.49 0.59 0.61 0.63 0.61 0.57 0.53 0.46 0.40 0.38 0.34 0.27 0.23 0.17 0.11 0.08 0.05 0.05
TIDAL CYCLE 40	120 130 140 150 160 170 180 193* 200 210 220 230 240 250 260 270 280 290 300 310 320	0.04 0.11 0.18 0.25 0.33 0.38 0.45 0.48 0.48 0.45 0.40 0.35 0.31 0.26 0.21 0.17 0.12 0.09 0.07

TEST 1-A

	Station	Corrected
		Conc.
TIDAL CYCLE 46	120 130 140 150 160 170 180 190 200* 210 220 230 240 250 260 270 280 290 310 320 340 350	0.04 0.08 0.14 0.20 0.25 0.28 0.31 0.33 0.34 0.33 0.32 0.30 0.28 0.26 0.22 0.19 0.15 0.12 0.09 0.06 0.05 0.04 0.03
TIDAL CYCLE 52	130 140 150 160 170 180 190 200 210* 220 230 240 250 260 270 280 290 300 310 320 330 340 350	0.01 0.03 0.05 0.07 0.11 0.14 0.18 0.21 0.21 0.19 0.18 0.16 0.15 0.13 0.12 0.11 0.09 0.07 0.05 0.04 0.03 0.02

	Station	CorrectedConc.
TIDAL CYCLE 58	150 160 170 180 190 200 210 220 230 240* 250 260 270 280 290 300 310 320 330 340 350	0.01 0.02 0.03 0.05 0.06 0.07 0.09 0.11 0.12 0.12 0.12 0.12 0.12 0.10 0.09 0.08 0.06 0.05 0.05 0.05

	Station	Corrected Conc.
TIDAL CYCLE 1	60* 70	11.06 1.80
TIDAL CYCLE 2	60 64.5* 70 80 90	6.58 7.05 5.98 1.96 0.31
TIDAL CYCLE 4	60 70 80* 90 100 110	2.62 4.06 4.68 4.03 2.32 0.16
TIDAL CYCLE 8	70 80 90 100 102.5* 110 120 130	0.96 1.81 3.01 3.32 3.37 3.16 2.21 0.52
TIDAL CYCLE 12	60 70 80 90 100 110 120* 130 140	0.09 0.18 0.28 0.43 0.85 1.75 2.41 1.80 1.24 1.09

	Station	Corrected Conc.	
TIDAL CYCLE 6	220 230 240 250 260 270 280 290 300* 310 320 330 350 350 370 380 390 410	0.01 0.03 0.63 0.93 1.22 1.50 1.79 1.83 1.79 1.50 1.50 1.50 1.26 0.79 0.40 0.13 0.09 0.07 0.03 0.01	
TIDAL CYCLE 24	220 230 240 250 260 270 280 290 310 320 340 350 350 360 370 380 390 410 420 430	0.09 0.09 0.09 0.13 0.18 0.27 0.36 0.45 0.54 0.59 0.68 0.63 0.36 0.39 0.32 0.29 0.18 0.09	

	Station	Corrected Conc.
TIDAL CYCLE 36	220 230 240 250 260 270 280 290 300 310 320 334* 340 350 360 370 380 410 420 430	0.02 0.04 0.06 0.10 0.14 0.20 0.24 0.32 0.37 0.53 0.57 0.53 0.53 0.24 0.18 0.16 0.14 0.12 0.10 0.08
TIDAL CYCLE 42	220 230 240 250 260 270 280 290 310 340* 350 340* 350 360 390 410 420 430	0.02 0.02 0.04 0.04 0.06 0.08 0.10 0.12 0.20 0.22 0.37 0.18 0.16 0.14 0.10 0.08 0.06 0.04

	Station	Corrected Conc.
TIDAL CYCLE 1	240 250 260 270 280 290 295* 300 310 320 330 340 350	0.03 0.18 0.44 1.12 2.41 4.84 5.22 4.84 1.65 0.18 0.03 0.01
TIDAL CYCIE 3	240 250 260 270 280 290 300* 310 320 330 340 350 370	0.19 0.68 1.34 1.88 2.47 3.12 3.44 3.07 2.24 1.05 0.36 0.18 0.10 0.06
TIDAL CYCLE 6	220 230 240 250 260 270 280 290 300 302.5* 310 320 330 340 350	0.07 0.14 0.21 0.53 0.96 1.86 1.96 1.97 1.86 1.74 1.29 0.89

		-
	Station	Corrected Conc.
TIDAL CYCLE 6 (cont'd)	360 370 380 390 400	0.56 0.14 0.09 0.04 0.03
(cont u)	410	0.01
TIDAL CYCLE 12	220 230 240 250 260 270 280 290 315* 320 330 350 370 360 370 410 420 430	0.04 0.07 0.12 0.26 0.40 0.90 1.48 1.47 1.32 1.10 0.59 0.49 0.31 0.21 0.13
TIDAL CYCLE 20	220 230 240 250 260 270 280 290 300 310 320* 330 340	0.02 0.03 0.09 0.14 0.22 0.31 0.41 0.65 0.84 0.91 0.95 0.86 0.81

	Station	Corrected Conc.
TIDAL CYCIE 20 (cont'd)	360 370 380 390 400 410 420 430	0.67 0.60 0.48 0.41 0.34 0.29 0.21
TIDAL CYCIE 30	220 230 240 250 260 270 280 290 300 322.5* 340 350 370 360 370 390 410 420 430	0.02 0.04 0.08 0.13 0.47 0.49 0.53 0.53 0.53 0.47 0.47 0.34 0.28 0.19 0.13
TIDAL CYCLE 40	220 230 240 250 260 270 280 290 300 310	0.02 0.02 0.03 0.04 0.07 0.10 0.18 0.30 0.32 0.32

	Station	Corrected Conc.
TIDAL CYCLE 40 (cont'd)	320 330* 340 350 360 370 380 390 400 410 420 430	0.34 0.34 0.34 0.32 0.30 0.26 0.22 0.18 0.14 0.10
TIDAL CYCLE 50	260 270 280 290 300 310 320 330* 340 350 360 370 380 390 410 420 430	0.04 0.08 0.12 0.17 0.19 0.21 0.21 0.21 0.14 0.10 0.08 0.06 0.04 0.04

The second

Sign of suppose

	Station	Corrected Conc.
TIDAL CYCLE 1	10 20 30 40 50 53.5* 60 70 80	0.09 0.19 0.52 2.87 11.66 17.40 5.57 0.26 0.09
TIDAL CYCLE 3	10 20 30 40 50 54* 60 70 80 90	0.34 0.59 1.50 2.40 5.92 4.46 1.16 0.22
TIDAL CYCLE 6	0 10 20 30 40 50 60* 79 80 90 100 110 120	0.24 0.49 0.73 1.26 2.16 2.75 3.03 2.82 2.07 1.59 1.13 0.73 0.31
TIDAL CYCLE 12	0 10 20 30 40 50 60 70 78.5* 80	0.25 0.43 0.60 0.85 1.40 1.74 2.06 2.32 2.43 2.41

-	Station	Corrected Conc.
TIDAL CYCLE 12 (cont'd)	90 100 110 120 130 140 150	2.08 1.65 1.21 0.79 0.38 0.10 0.04
TIDAL CYCLE 20	20 30 40 50 60 70 80 90* 100 110 120 130 140 150 160 170 180 190 200	0.35 0.48 0.66 0.83 1.06 1.44 1.82 1.96 1.79 1.79 1.04 0.75 0.48 0.35 0.18 0.14 0.10 0.07
TIDAL CYCLE 30	0 10 20 30 40 50 60 70 80 90 98* 100 110 120 130 140 150 160 170 180	0.19 0.20 0.23 0.31 0.40 0.54 0.79 1.04 1.27 1.47 1.52 1.50 1.36 1.15 1.01 0.85 0.65 0.47 0.36 0.28

	Station	Corrected Conc.
TIDAL CYCLE 30 (cont'd)	190 200 210 220 230 240 250	0.23 0.19 0.16 0.14 0.12 0.11 0.08
TIDAL CYCLE 40	0 10 20 30 40 50 60 70 80 90 120* 130 140 150 150 160 190 210 250 260 270 280 290 300	0.10 0.15 0.15 0.15 0.56 0.56 0.57 0.64 0.64 0.65 0.65 0.65 0.65 0.65 0.65 0.65 0.65

	Station	Corrected Conc.
TIDAI. CYCLE 1	80 90 100 110 116.75* 120 130 140	0.05 0.15 0.23 4.41 9.71 9.04 1.14 0.05
TIDAL CYCLE 3	70 80 90 100 110 125* 130 140 150	0.08 0.12 0.17 0.28 1.70 4.95 5.45 5.15 2.09 0.45 0.13
467		
TIDAL CYCLE 6	80 90 100 110 120 130 139.5* 140 150 160 170 180 190 200	0.03 0.10 0.18 0.49 1.04 2.05 3.25 3.24 2.39 1.26 1.00 0.39 0.13 0.07
TIDAL CYCLE 12	80 90 100 110 120 130 140 150	0.01 0.02 0.04 0.09 0.36 0.77 1.33 1.86 2.09

	Station	Corrected Conc.
TIDAL CYCLE 12 (cont'd)	170 180 190 200 210 220 230 240 250 260	1.80 1.33 0.94 0.61 0.39 0.38 0.36 0.23 0.12 0.07 0.06
TIDAL CYCLE 20	100 110 120 130 140 150 160 170 175* 180 190 200 210 220 230 240 250 260 270 280 290 300 310	0.02 0.03 0.08 0.23 0.42 0.62 0.90 1.05 1.05 1.01 0.94 0.86 0.84 0.81 0.70 0.56 0.41 0.25 0.16 0.14 0.08
TIDAL CYCLE 30	110 120 130 140 150 160 170 180 190 200	0.02 0.05 0.08 0.11 0.16 0.26 0.36 0.43 0.49 0.56

	Station	Corrected Conc.
TIDAL CYCLE 30 (cont'd)	220 230* 240 250 260 270 280 290 300 310 320	0.62 0.64 0.57 0.52 0.46 0.43 0.38 0.33 0.28 0.16 0.08

	Station	Corrected Conc.
TIDAL CYCLE 1	90 100 110 116.5* 120 130 140	0.06 0.17 4.38 10.43 9.12 0.81 0.08
TIDAL CYCLE 3	70 80 90 100 110 120 124.5* 130 140 150 160 170 180	0.01 0.04 0.08 0.19 1.37 4.13 4.58 4.10 2.18 0.55 0.18 0.17 0.13
TIDAL CYCLE 6	70 80 90 100 110 120 137* 140 150 160 170 180 190 200 210	0.05 0.07 0.08 0.16 0.26 1.11 2.57 2.83 2.78 2.26 1.17 0.60 0.30 0.16 0.07 0.04

	Station	Corrected Conc.
TIDAL CYCLE 12	90 100 110 120 130 140 150 158.5* 160 170 180 190 200 210 220 230 240 250 260 270	0.03 0.06 0.12 0.30 0.59 0.94 1.29 1.42 1.41 1.10 0.88 0.73 0.57 0.42 0.09 0.06 0.04 0.03
TIDAL CYCLE 20	90 100 110 120 130 140 150 160 170 180 190* 200 210 220 230 240 250 260 270 280 290 300 310	0.02 0.03 0.05 0.09 0.12 0.17 0.31 0.47 0.59 0.73 0.76 0.73 0.62 0.55 0.47 0.44 0.44 0.45 0.23 0.17

	Station	Corrected Conc.
TIDAI, CYCIE 30	120 130 140 150 160 170 180 290 210 220 230* 240 250 260 270 280 290 300 310 320 330 340 350	0.02 0.05 0.10 0.15 0.33 0.33 0.346 0.54 0.54 0.46 0.41 0.30 0.13 0.11 0.08 0.05 0.03
TIDAL CYCLE 40	130 140 150 160 170 180 190 200 210 220 240 250* 26c 270 280 290 300 310 320 330	0.03 0.05 0.07 0.10 0.14 0.17 0.21 0.26 0.31 0.32 0.34 0.36 0.38 0.36 0.32 0.29 0.26 0.22 0.19 0.15

	Station	Corrected Conc.
TIDAL CYCLE 40 (cont'd)	340 350 360 370	0.10 0.09 0.07 0.07
TIDAL CYCLE 50	150 160 170 180 190 200 210 220 240 250 260 270* 280 290 300 310 320 340 350 370	0.02 0.03 0.09 0.12 0.15 0.17 0.17 0.19 0.21 0.22 0.24 0.29 0.31 0.28 0.26 0.22 0.21 0.19 0.19 0.19

	Station	Corrected Conc.
TIDAL CYCIE 1	80 90 100 110 116.50* 120 130 140	0.02 0.06 0.11 7.60 11.52 11.02 0.49 0.14
TIDAL CYCLE 3	70 80 90 100 110 120 126.5* 130 140 150	0.06 0.09 0.13 0.19 1.57 4.86 5.81 5.38 1.60 0.33
TIDAL CYCLE 6	70 80 90 100 110 120 130 139.5* 140 150 160 170 180 190 200 210	0.01 0.05 0.10 0.18 0.26 1.20 2.51 5.34 3.32 2.30 1.26 0.68 0.35 0.18 0.13

	Station	Corrected Conc.
TIDAL CYCLE 12	90 100 110 120 130 140 150 158* 160 170 180 190 200 210 220	0.02 0.03 0.06 0.33 0.70 0.91 1.38 1.75 1.71 1.39 1.10 0.84 0.61 0.36 0.25 0.17
TIDAL CYCLE 20	110 120 130 140 150 160 170 180 197* 200 210 220 230 240 250 260 270 280 290 310	0.05 0.08 0.12 0.39 0.55 0.70 0.84 0.81 0.55 0.22 0.17 0.12

Station		Corrected
		Conc.
TIDAL CYCLE 30	130 140 150 160 170 180 190 200 210 220 230 232.5* 240 250 260 270 280 290 300 310 320 330 340 350	0.05 0.07 0.10 0.15 0.20 0.26 0.31 0.45 0.48 0.48 0.48 0.46 0.41 0.38 0.33 0.28 0.25 0.20 0.15 0.07
TIDAL CYCLE 40	130 140 150 160 170 180 190 200 210 220 230 240 250* 260 270 280 290 300 310	0.03 0.05 0.05 0.07 0.09 0.14 0.19 0.24 0.29 0.32 0.34 0.38 0.38 0.38 0.38 0.38 0.31 0.27 0.27

	Station	Corrected Conc.
TIDAL CYCLE 40 (cont'd)	320 330 340 350 360 370	0.15 0.14 0.14 0.12 0.10 0.09
TIDAL CYCLE 50	150 160 170 180 190 200 210 220 230 240 250 260 270* 280 290 300 310 320 340 350 360 370	0.03 0.05 0.06 0.07 0.08 0.09 0.12 0.15 0.17 0.24 0.21 0.24 0.21 0.17 0.14 0.12 0.10 0.09 0.09

APPENDIX IV

- A. Auxiliary Data on Salinity Distribution, High and Low Water Profiles, and Fresh Water Inflow.
- B. Table of Cross Sectional Area, Intersectional Volume, and Accumulated Volume for each 5000 ft. Station.

DELAWARE RIVER MODEL Salinity Profile Data Tests 2-3-4

Bottom	27,270 22,801 17,302 10,656 7,963 5,099 3,495 2,640 2,640
Test 1, Mean Tide Low flow (5,905) Top Middepth	27,155 21,999 16,958 10,541 6,588 4,640 3,437 2,040 825 665 183 92
Test ? Low f.	26,697 21,999 15,812 9,625 6,588 1,010 3,094 1,971 1,971 1,971
de 000) Bottom	26,812 23,947 19,135 11,114 8,114 8,114 894 711 711 255 57 255 34
; 3, Mean Tide Iflow (49,000) Middepth	26,12h 23,145 17,302 9,854 6,784 1,994 1,994 61.9 61.9 61.9 57 61.9 57 34
Test High Top	23,260 21,197 13,979 6,875 1,169 1,169 1,169 1,169 1,13 206 69 34
le DO) Bottom	27,385 25,666 25,731 15,182 2,182 3,553 3,553 3,553 172 80 80 50
2, Mean Tide flow (20,200) Middepth	27,385 26,208 24,291 18,677 14,265 9,453 5,226 1,902 1,902 80 332 1,72 80 50 50
Test Mean Top	27,385 25,208 23,603 15,583 12,030 7,734 1,813 2,854 2,854 1,60 69 1,60 69
1000 ft Channel Stations	430 430 350 350 250 250 150 150 120 120 120

Quantities are PPM total salt. Subtract average fresh-water salinity (40 PPM) and divide result by 1.81 to obtain equivalent chlorine. NOTE:

DELAWARE RIVER MODEL Salinity Profile Data Tests 5-6-7

(b) Bottom	13,291 6,646 3,953 1,054 504 69
7, Neap Tide flow (20,200) Middepth	12,375 6,416 3,724 1,054 390 69
Test Nean f	11,771 5,328 2,635 2,635 894 321 69
ide O) Bottom	8,250 3,667 2,200 1,375 929
6, Spring Tide I flow (20,200) Middepth B	7,906 3,667 2,040 1,375 92
Test Mean Top	6,989 2,521 1,306 1,215 206 92
3 . 3ottom	9,854 4,469 2,842 504 275 115
5, Mean Tide flow (20,200) Middepth	8, 823 4, 469 2, 682 504 252 115
Test Mean Top	7,9390 1,437 1,834 527 229 11.5
1000 ft Channel Stations	250 210 190 160 150 130

Quantities are PPM total salt. Subtract average fresh-water salinity (40 PPM) and divide result by 1.81 to obtain equivalent chlorine. NOTE:

DELAMARE RIVER MCDEL Data For High and Low Vater Profile

4		
	Low Mater Elev	င္ဝင္ဝင္ဝင္ဝင္ဝင္ဝင္ဝင္ဝင္ ဆမ္းႏွစ္တဲ့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တို့တင္တင္တင္တင္တင္တင္တင္တင္တင္တင္တင္တင္တင္တ
Meap	High Water Elev	www.www.woogw. wwocregocoddy.woogw
Spring	Low Water Elev	
Spr	High Water Blev	
	Low Vater Elev	
Mean	High Vater Elev	7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7
	1000 ft Channel Station	127 233 333 333 150 150 150 150 150 150 150 150 150 150
Location	Gage	Miah Maull Ship John Yoodland Beach Artificial Island Reedy Point New Castle Edge Moor Marcus Hook Faldwin Fort Mifflin Philadelphia Torresdale Rurlington Florence Trenton

FRESH-WATER INFLOW DISTRIBUTION

Inflow Location	Prototype C.F.S. Mean Flow Tests 1-A,1,2,5,6,7	Prototype C.F.S. High Flow Test 3	Prototype C.F.S. Low Flow Test 4	Remarks
Trenton Rancocas	12,350 8 7 5	31,300 0	3,000 0	Combined with Schuylkill
Schuylkill Christina	3,250 1,100	8,700 2,650	2,000 480 0	Combined with
Salem	450 425	1,100	0	Christina Combined with
Maurice	1,750	4,250	425	Christina
(Total Fresh-wate	er)) 20,200	49,000	5,905	

CROSS-SECTIONAL AREAS AND SEGMENT VOLUMES - DELAWARE RIVER

Station	Cross-section Area x 10 ⁻² ft ²			Area Volume			j	Accumulated Volume x 10 ⁻⁶ ft ³			
	low tide	mean tide	high tide	low tide	mean tide	high tide	low tide	mean tide	high tide		
- 160 - 155 - 150 - 145 - 140 - 135 - 130 - 125 - 120 - 115 - 110 - 105 - 100 - 95 - 90 - 85 - 80 - 75 - 70	22 70 117 228 178 156 184 273 212 212 192 184 214 209 242 212 239 223 289	53 104 148 298 231 203 226 376 253 223 272 232 264 251 315 261 289 287 349	83 138 178 367 284 250 268 479 294 233 351 279 313 292 388 310 338 350 408 456	23 47 86 102 84 85 114 121 106 101 94 100 106 113 113 113 113 114 128 147	39 63 111 133 109 107 150 157 128 133 126 124 128 142 144 138 144 159 182	55 79 136 163 134 129 186 195 150 164 158 148 151 170 174 162 172 189 216	0 23 70 156 258 342 427 541 662 768 869 963 1063 1169 1282 1395 1508 1624 1752 1899	39 102 213 346 455 562 712 869 997 1129 1255 1379 1508 1649 1793 1930 2074 2233 2414	0 55 134 270 433 567 696 882 1075 1225 1389 1547 1695 1846 2016 2190 2352 2524 2713 2929		
- 65 - 60	23 ⁴	377 306	378	133	171 167	209		2585	3138		

Station	Cross-section Area x 10 ⁻² ft ²			Segment Volume x 10 ⁻⁶ ft ³			Acc 		
	low tide	mean tide	high tide	low tide	mean tide	high tide	low tide	mean tide	high tide
- 55	278	360	441			701	2160	2752	3343
- 45	303	366	428	229	277	324	2550	3232	3913
- 40	373	474	575	169	213	257	2719	3445	4170
- 35	392	488	583	191	241	290	2910	3685	4460
- 30	348	459	569	185	237	288	3095	3922	4748
- 25	387	520	753	184	245	306	3279	4167	5054
- 20	482	594	705	217	279	34ž	3496	4446	5396
- 15	390	527	664	218	280	342	3714	4726	5738
- 10	415	511	606	201	259	317	3915	4985	6055
- 05	440	530	620	214	260	306	4129	5425	6361
00	585	733	880	256	316	375	4385	5561	6736
+ 05	649	761	873	308	373	438	4693	5934	7174
+ 10	654	746	837	326	377	428	5019	6311	7602
+ 15	543	603	662	299	337	375	5318	6648	7977
+ 20	740	817	894	321	355	389	5639	7003	8366
+ 25	604	677	749	336	374	411	5975	7376	877?
+ 30	635	708	781	310	347	383	6285	7723	9160
+ 35	601	673	744	309	346	382	6594	8068	9542
+ 40	852	1001	1149	363	418	473	6957	8486	10015
+ 45	701	819	937	357	75 7	490	7314	8909	10505
+ 50	791	901	1010	373	430	487	7697	933 9	10992
+ 55	1010	1173	1335	450	518	586	8137	9858	11578
				507	590	673			

Station	Cross-section Area x 10 ⁻² ft ²			177	egment clume 0-6 ft	3	Accumulated Volume x 10-6 ft ³		
	low tide	mean tide	high tide	low tide	mean tide	high tide	low tide	mean tide	high tide
+ 60 + 65 + 70 + 75 + 80 + 85	1016 899 1175 796 754 885	1185 1059 1289 887 937 1089	1354 1218 1402 978 1119 1292	479 518 493 388 412 438	561 586 546 457 509 531	643 654 598 525 605 624	8644 9123 9641 10134 10522	1009 11595 12140 12597 13105	12251 12894 13548 14146 14671 15276
+ 90 + 95 + 100 + 105 + 110 + 115 + 120 + 125 + 130 + 135 + 140 + 145	866 958 1008 846 991 1047 1016 1378 1074 1219 1375 1230	1039 1150 1232 1047 1157 1174 1218 1547 1250 1408 1589 1444	1212 1341 1456 1248 1322 1301 1420 1716 1425 1597 1803 1658	456 491 463 459 509 516 598 613 573 649 651 630	547 595 570 551 578 598 691 699 665 750 758 734	638 699 676 642 646 680 783 785 756 851 865 838	11372 11828 12319 12782 13241 13750 14266 14864 15477 16050 16699 17350	19878 20636	22205 23056 23921
+ 150 + 155 + 160 + 165	1292 1292 1356 1292	1494 1506 1585 1549	1696 1720 1814 1805	646 662 662 821	750 773 784 945	854 883 905 1068	17980 18626 19288 19950	21369 22119 22892 23676	24759 25613 26496 27501

Station	Cross-section Area x 10-2 ft ²			Segment Volume x 10-6 ft ³			Accumulated Volume x 10 ⁻⁶ ft ³		
	low tide	mean tide	high tide	low tide	mean tide	<u>high</u> tide	low tide	mean tide	high tide
+ 170	1993	2231	2469	843	954	1065	20771	24620	28469
+ 175	1381	1587	1793	646	739	833	21614	25574	29534
+ 180 + 185	1202	1371 1845	1540 2083	702	804	906	22260	26314 27118	30367 31273
+ 190	1673	1908	2142	820	939	1057	23782	28056	32330
+ 195	1687	1923	2159	840 851	958 973	1075	24622	29014	33405
+ 200	1717	1968	2219	832	966	1099	25473	29986	34499
+ 205 + 210	1609 1687	1892 1970	2175	824	965	1107	26305 27129	30952 31917	35598 36705
+ 215	1681	2058	2435	842	1007	1172	27971	3585 ji	37977
+ 220	1559	1902	5544	810 817	990 1003	1170 1189	28781	33914	39047
+ 225	1709	2111	2513	948	1152	1355	29598	34917	40236
+ 230 + 235	2082 2154	2493 2649	2904 3144	1059	1286	1512	30546 31605	36069 37354	41591 43103
+ 240	1832	2138	2444 2144	997	1198	1398	32602	38552	44501
+ 245	1776	2120	2464	902 888	1065	1227	33504	39616	1,5728
+ 250	1776	2123	2469	1002	1180	1358	34 3 92	40677	46961
+ 255 + 260	2232 2174	2598 2517	2964 2860	1102	1279	1457	35394 36496	41857 43136	48319 49776
+ 265	2413	2763	3112	1147	1320	1493	37643	44456	51269
+ 270	22 ⁴ ⁴	2650	3055	1164	1353 1384	1542 1572	38807	45809	52811
+ 275	2536	2884	3232	1308	1497	1685	40002	47193	54383

Station	Cross-section Area x 10 ⁻² ft ²			Segment Volume x 10-6 ft ³			Accumulated Volume x 10 ⁻⁶ ft ³			
	low tide	mean wide	high tide	low tide	mean tide	high tide	low tide	mean tide	high tide	
+ 280 + 285 + 290 + 295 + 300 + 305 + 310 + 315 + 320 + 325 + 330 + 335 + 340 + 345 + 350 + 355 + 360 + 365 + 370		3101 3237 3439 3624 4074 4137 4328 4381 4744 5090 4790 4472 4895 6118 6357 6302 6600 7168 7675		1358 912 1486 1605 1707 1763 1833 1930 2073 2100 1976 1978 2310 2607 2658 2739 2934 3155 3551	1584 1169 1766 1924 2053 2117 2178 2282 2458 2470 2316 2342 2753 3119 3165 3225 3442 3711 4225		41310 42668 43580 45066 46671 48378 50141 51974 53904 55977 58077 60053 62031 64341 66948 69606 72345 75279 78434	48689 50274 51443 53208 55132 57185 59301 61479 63760 66219 68688 71005 73546 76099 79218 82372 85607 89049 92759	56068 57879 59305 61350 63593 65991 68461 70983 73616 76460 79300 81956 84661 87857 91487 95158 98869 102819 107085	
+ 375 + 380	7694 8729	9228 10456	10762	4104 4728	4919 5535	5735 6342	81935 86089	96985 101904	111984	

Station		oss-sec Area : 10 ⁻²		¥o.	gment lume -6 ft ³		Ac	d +3	
	low tide	mean tide	high tide	low tide	mean tide	high tide	low tide	mean tide	high tide
+ 385	8573	10219	11865	4365	5129	5894	90817	107439	124061
+ 390	10316	12145	13974				95182	112569	129955
+ 395	10455	12463	14470	5193	6152	7111	100375	118721	137066
+ 400	1057 7	12674	14770	5258	6284	73.10	105633	125005	144376
+ 405	11468	13719	15969	6326	7523	8719	111959	132527	153095
+ 410	12131	14391	16650	5849	6928	8006	117808	139455	161101
+ 415	12927	15156	17385	6265	7387	8509	124073	146824	169610
-	•		1	6636	7765	8895			-
+ 420	13617	15907	18196	6960	8112	9264	130709	154607	178505
+ 425	14224	16543	18862	7638	8960	10282	137669	162719	187769
+ 430	16328	1.9296	22264		0,000	-0-0-	145307	171679	198051

DISTRIBUTION LIST

Copie	s Addressee	<u>Copies</u> <u>Addressee</u>	
1	Chief of Naval Operations Navy Department Washington 25, D. C. Attn: Op=533D	1 Office of Naval Research Branch Office 1030 East Green Street Pasadena 1, California	
1	Chief of Naval Research Navy Department Washington 25, D. C. Attn: Code 466	 Office of Naval Research Branch Office 1000 Geary Street San Francisco, California 	
2	Geophysics Branch Code 416 Office of Naval Research Washington 25, D. C.	6 Director Naval Research Laboratory Washington 25, D. C. Attn: Technical Information Officer	
1	Office of Naval Research Contract Administrator Southeastern Area c/o George Washington University 2110 G Street, N. W.	8 U. S. Navy Hydrographic Offic Washington 25, D. C. Attn: Division of Oceanographic	
1	Washington 7, D. C. Mr. W. B. Girkin	2 Assistant Naval Attache for Roamerican Embassy Navy 100	esearch
	ONR Resident Representative Institute for Cooperative Research	Fleet Post Office, New York	
	1315 St. Paul Street Baltimore 2, Maryland	 British Joint Services Mission Main Navy Building Washington 25, D. C. 	1
2	Officer-in-Charge Office of Naval Research London Branch Office Navy Number 100 Fleet Post Office New York, N. Y.	 Chief, Bureau of Ships Navy Department Washington 25, D. C. Attn: Code 847 	
1	Office of Naval Research Branch Office 346 Broadway	 Chief, Bureau of Yards and D Navy Department Washington 25, D. C. 	ocks
1	New York 13, N. Y. Office of Naval Resea: h Branch Office Tenth Floor, The John Crerar Library Building	 Director, U. S. Navy Electron Laboratory San Diego 52, California Attn: Codes 550, 552 	nics

86 East Kandolph Street

Chicago, Illinois

Copies Addressee Copies Addressee Commander, Naval Ordnance Laboratory 1 Commanding General White Oak Research and Development Division Silver Spring, Maryland Department of the Army Washington 25, D. C. Amphibious Training Command 1 Naval Amphibious Base U. S. Army Beach Erosion Board 1 Little Creek, Virginia 5201 Little Falls Road, N. W. Washington 16, D. C. Project Arowa U. S. Naval Air Station District Engineer Building R-48 Baltimore District Norfolk, Virginia Corps of Engineers, U. S. Army P.O. Box 1715 Department of Aerology Baltimore 3, Maryland U. S. Naval Postgraduate School Monterey, California 1 Waterways Experiment Station U. S. Army Engineers 1 U. S. Naval Underwater Sound Vicksburg, Mississippi Attn: Mr. Alan G. Skelton, Head Laboratory New London, Connecticut Research Center Library U. S. Navy Mine Counter Measure District Engineers Office Station U. S. Army Engineers Panama City, Florida Norfolk, Virginia Commanding Officer Cambridge Field Station 5 Armed Services Technical Information 230 Albany Street Center Cambridge 39, Massachusetts Documents Service Center Knott Building Dayton 2, Ohio 1 Director, U. S. Coast and 1 Assistant Secretary of Defense for Geodetic Survey Research and Development Department of Commerce Pentagon Building Washington 25, D. C. Washington 25, D. C. Attn: Committee on Geophysics 1 Commandant (OAC) U. S. Coast Guard and Geography Washington 25, D. C. Commanding General Research and Development Division 2 U. S. Fish and Wildlife Service Department of the Air Force Department of the Interior Washington 25, D. C. Washington 25, D. C. Attn: Dr. L. A. Walford

Copies Addressee

- U. S. Fish and Wildlife Service
 450 B Jordan Hall
 Stanford University
 Stanford California
- U. S. Fish and Wildlife Service South Atlantic Fishery Investigations
 P. O. Box 283
 Brunswick, Georgia
- U. S. Fish and Wildlife Service
 P. O. Box 3830
 Honolulu, T. H.
- U. S. Fish and Wildlife Service Chesapeake Shellfish Investigations
 P. O. Box 151 Annapolis, Maryland Attn: Mr. J. B. Engle
- Mr. Paul F. Springer, Biologist U. S. Department of Interior Fish and Wildlife Service Patuxent Research Refuge Laurel, Maryland
- U. S. Fish and Wildlife Service Woods Hole Massachusetts
- U. S. Fish and Wildlife Service Fort Crockett
 Galveston, Texas
- U. S. Fishery Laboratory Beaufort, North Carolina
- Librarian
 U. S. Geological Survey
 Room 1033
 General Services Adm. Building
 Washington 25, D. C.

Copies Addre ce

- National Research Council
 2101 Constitution Avenue
 Washington 25, D. C.
 Attn: Committee on Undersea
 Warfare
- Dr. Garbis H. Keulegan Hydraulics Division National Eureau of Standards Washington 25, D. C.
- Division of Water Pollution Control U. S. Public Health Service Federal Security Agency Washington 25, D. C. Attn: Mr. A. F. Bortsch
- Office of Technical Services
 Department of Commerce
 Washington 25, D. C.
- Weather Eureau Library
 24th and M Streets, N. W.
 Washington 25, D. C.
- Department of Research and
 Education
 State of Maryland
 Solomons, Maryland
 Attn: Dr. R. V. Truitt, Director
- Director
 Maryland Department of Geology,
 Mines and Water Resources
 c/o The Johns Hopkins University
 Baltimore 18, Maryland
- 2 Eoard of Natural Resources Dept. of Tidewater Fisheries State Office Building Annapolis, Maryland Attn: Mr. Arthur H. Brice, Chairman

Copies Addressee

- 1 Mr. Edwin M. Barry
 Chief, Inland Fish Management
 State Game and Inland Fish Commission
 516 Munsey Building
 Baltimore 2, Maryland
- Water Pollution Control Commission
 State of Maryland
 2203 North Charles Street
 Baltimore 18, Maryland
- Department of Information State Office Building Annapolis, Maryland
- Director
 Virginia Fisheries Laboratory
 Gloucester Point, Virginia
- Director
 Division of Statutory Research
 and Drasting
 State Capitol
 Richmond 19, Virginia
- Municipal Reference Library
 2230 Municipal Building
 New York 7, N. Y.
 Attn: Miss Amelia H. Munson
- Baltimore County Health DepartmentTowson 4, MarylandAttn: Mr. V. R. Sullivan
- 1 Talbot County Free Library
 Easton, Maryland
 Attn: Miss Sarah F. Cockey
 Librarian
- Bingham Oceanographic FoundationYale UniversityNew Haven, Connecticut

- 1 Dr. R. B. Montgomery
 Professor of Oceanography
 Brown University
 Providence, R. I.
- Department of Conservation
 Cornell University
 Ithaca, New York
 Attn: Dr. J. Ayers
- 1 The Oceanographic Institute Florida State University Tallahassee, Florida
- l Director
 Narragansett Marine Laboratory
 Kingston, Rhode Island
- Department of Meteorology and Oceanography
 New York University
 New York, N. Y.
- 1 Marine Physical Laboratory
 Scripps Institution of Oceanography
 Point Loma, California
- Director
 Scripps Institution of Oceanography
 La Jolla, California
- 2 Head, Department of Oceanography Texas A and M College College Station, Texas
- Director
 Marine Laboratory
 University of Miami
 Coral Gables, Florida
- 1 Allen Hancock Foundation University of Southern California Los Angeles 7, California

Copies Addressee 2 Department of Oceanography University of Washington Seattle 5, Washington

Director
 Woods Hole Oceanographic Institution
 Woods Hole, Massachusetts

Attn: Dr. Richard H. Fleming

Bears Bluff Laboratories Wadmalaw Island, S. C. Attn: Mr. G. Robert Lunz, Director

Librarian

- Director, Hawaii Marine Laboratory
 University of Hawaii
 Honolulu, T. H.
- Institute of Fisheries Research
 University of North Carolina
 Morehead City, North Carolina
- Director
 Institute of Marine Science
 Port Aransas, Texas
- Wm. G. Kerckhoff Laboratories of the Biological Sciences California Institute of Technology Corona del Mar, California
- 1 Marine Biological Laboratory Woods Hole, Massachusetts Attn: Librarian
- Marine LaboratoryDuke UniversityBeaufort, North Carolina
- Miss Jessie Bell MacKenzie Librarian Museum of Comparative Zoology Harvard University Cambridge, Massachusetts

- Stone Biological Laboratory
 Put-in-Bay
 Ohio
- Department of Zoology
 Rutgers University
 New Brunswick, New Jersey
 Attn: Dr. H. H. Haskin
- Brown University Providence, Rhode Island Attn: Dr. George W. Morgan Applied Math. Dept.
- Institute for Cooperative Research
 The Johns Hopkins University
 1315 St. Paul Street
 Baltimore 2, Maryland
 Attn: Librarian
- 1 The Johns Hopkins University
 Baltimore 18, Maryland
 Attn: Librarian (Acquisitions Dept.)
- Department of Sanitary Engineering
 The Johns Hopkins University
 Baltimore 18, Maryland
 Attn: Dr. Abel Wolman (1)
 "Dept. Library (1)
- Department of Civil Engineering
 The Johns Hopkins University
 Baltimore 18, Maryland
 Attn: Dr. George S. Benton
- The Isaiash Bowman School of Geography
 The Johns Hopkins University Baltimore 18, Maryland Attn: Dr. George Carter

Copies Addressee

4 University of Maryland College Park, Maryland

Attn: Miss Virginia Phillips
Ass't Reference Librarian

- " Librarian, Maryland Division
- " Department of Biology
- " Bureau of Public Administration Christian L. Larsen, Director
- Department of Engineering University of California Berkeley, California
- University of California
 Department of Meteorology
 405 Hilgard
 Los Angeles 24, California
- Director
 Lamont Geological Observatory
 Columbia University
 Torrey Cliff
 Palisades, New York
- Department of Zoology
 Oklahoma A and M College
 Stillwater, Oklahoma
 Attn: Dr. I. Eugene Wallen
- Limnology Library
 Biology Building
 University of Wisconsin
 Madison 5, Wisconsin
- 1 Enoch Pratt Free Library
 400 Cathedral Street
 Baltimore 1, Maryland
 Attn: Miss Elizabeth C. Litsinger
 Head, Maryland Department

- The Oyster Institute of North
 America
 6 Mayo Avenue, Bay Ridge
 Annapolis, Maryland
 Attn: Mr. David H. Wallace,
 Director
- 1 Atlantic States Marine Fisheries 22 West First Street Mount Vernon, New York Attn: Mr. Wayne D. Heydecker
- California Academy of Sciences
 Golden Gate Park
 San Francisco, California
 Attn: Dr. R. C. Miller
- Librarian
 California Fisheries Laboratory
 Terminal Island Station
 San Pedro, California
- The New York Public Library Astor, Lenox and Tilden Foundations, Acquisitions Division Fifth Avenue and 42nd Street New York 18, N. Y.
- Department of Zoology
 University of Cape Town
 Rondebosch
 SOUTH AFRICA
 Attn: Dr. N. A. Millard (1)
 '' Dr. J. H. Day (1)
- 1 Fisheries Division C.S.I.R.O.
 P. O. Box 21
 Cronulla, N. S. W.
 AUSTRALIA
 Attn: Dr. D. J. Rochford

Copies Addressee Atlantic Biological Station Fisheries Research Board of Canada St. Andrews New Brunswick, Canada Dr. John P. Tully Pacific Oceanographic Group c/o Pacific Biological Station Nanaimo

Pacific Biological Station
 Fisheries Research Board of Canada
 Nanaimo
 British Columbia, CANADA

British Columbia, CANADA

- Naval Research Establishment c/o Fleet Mail Office Halifax Nova Scotia, CANADA
- Fisheries Research Board of Canada
 Zoological Building
 University of Toronto
 Toronto, Ontario
 CANADA
- Institute of Oceanography
 University of British Columbia
 Vancouver, Canada
 Attn: Dr. W. M. Cameron (1)
 '' Librarian (1)
- Tidal and Current Survey Canadian Hydrographic Service Dept. of Mines and Technical Surveys Ottawa, CANADA
- Conseil International pour l'Exploration de la Mer Bibliotheque Postbox 20 Charlottenlund, Denmark Attn: Librarian
- Liverpool Observatory and Tidal Institute Bidston, Birkenhead, England Attn: Librarian

- Dr. M. S. Longuet-Higgins Cambridge University Cambridge, England
- 1 Marine Biological Association of the United Kingdom
 The Plymouth Laboratory
 Citadel Hill
 Plymouth, England
- Dr. J. R. Lumby, Editor "Journal du Conseil" Fisheries Laboratory Lowestoft Suffolk, England
- 1 Librarian National Institute of Oceanography Wormley, near Godalming Surrey ENGLAND
- 1 Dr. J. Proudman
 Professor of Oceanography
 University of Liverpool
 Liverpool, England
- L'Ingenieur Hydrographe en Chef Gougenheim Comite Central d'Oceanographie et d'Etude des Cotes 13 Rue de l'Universite Paris VII FRANCE
- Deutsches Hydrographisches
 Institut
 Seewartenstrasse 9
 Hamburg 11
 GERMANY
 Attn: Dr. G. Bohnecke, President
- 1 Dr. S. Jones
 Central Fisheries
 Barrackpore P. O.
 via: Calcutta
 INDIA

Copies Addressee Food and Agriculture Organization of the United Nations Viale delle Terme di Caracalla Rome ITALY Attn: Miss Kristine Lomsdal, Librarian Dr. Koji Hidaka Geophysical Institute Tokyo University Tokyo, Japan Masao Kametaka Librarian Faculty of Agriculture Tohoku University Sendai, Japan Dr. Kanji Suda Chief Hydrographer Hydrographic Office 5 - Tsukiji, Chuo-Ku Tokyo **JAPAN** 1 International Hydrographic Bureau Monte Carlo MONACO 1 Musee Oceanographique de Monaco Monaco-Ville MONACO Institut Français d'Oceanie Boite Postale No. 4 Noumea NEW CALEDONIA Dr. E. I. Robertson Department of Geophysics D. S. I. K. Wellington

NEW ZEALAND

- 1 Geologisch Instituut
 Rijks-University
 Melkweg 1
 Groningen,
 THE NETHERLANDS
- 1 Dr. P. Groen
 Kon. Ned. Meteorologisch
 Instituut Oceanografische
 Afdeling, de Bilt
 THE NETHERLANDS
- Universitetet OsloBlindern, OsloNorwayAttn: Librarian
- Geofysisk InstituttAllegaten 70BergenNorway
- Professor H. U. Sverdrup, Dir.
 Norsk Polarinstitutt
 Observatoreight 1
 Oslo
 Norway
- Norwegian Embassy 3401 Massachusetts Ave., N. W. Washington, D. C. Attn: Fisheries Attache
- 1 Marine Station
 Millport, Isle of Cumbrae
 SCOTLAND
 Attn: Dr. H. Barnes
- 1 Dr. Hans Pettersson Oceanografiska Institutet Box 1038, Goteborg 4 SWEDEN
- Library
 Kungl. Fiskeristyrelsen med
 Statens, Fiskeriforsok
 Hydrographic Dept.
 Box 1038, Goteborg 4
 SWEDEN

- Sea Fisheries Research StationHaifa P. O. Box 699Israel
- Fiskeridirektøren
 Biblioteket
 Postboks 226
 Bergen, Norway
 Margit Monstad, Librarian
- Director
 Bermuda Biological Station
 for Research
 St. George's, Bermuda
- Project Officer
 Laboratory of Oceanography
 Woods Hole, Massachusetts
- 1 Dr. F. Møller Norwegian Defense Research Institute Akershus, Oslo, Norway
- Mr. John B. Glude, Chief
 Clam Investigations
 U. S. Dept. of the Interior
 Fish and Wildlife Service
 Boothbay Harbor, Maine